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(54) Title: SEED TRAIT GENES

(57) Abstract: Recombinant polynucleotides and methods for modifying the phenotype of a plant are provided. In particular, the phenotype that is being modified is a plant's seed characteristics.

SEED TRAIT GENES**RELATED APPLICATION INFORMATION**

The present invention claims the benefit from US Provisional Patent Application Serial
5 Nos. 60/166,228 filed November 17, 1999 and 60/197,899 filed April 17, 2000 and "Plant Trait
Modification III" filed August 22, 2000.

FIELD OF THE INVENTION

This invention relates to the field of plant biology. More particularly, the present invention pertains to compositions and methods for phenotypically modifying a plant.

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BACKGROUND OF THE INVENTION

Transcription factors can modulate gene expression, either increasing or decreasing (inducing or repressing) the rate of transcription. This modulation results in differential levels of gene expression at various developmental stages, in different tissues and cell types, and in response to different exogenous (e.g., environmental) and endogenous stimuli throughout the life cycle of the organism.

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Because transcription factors are key controlling elements of biological pathways, altering the expression levels of one or more transcription factors can change entire biological pathways in an organism. For example, manipulation of the levels of selected transcription factors may result in increased expression of economically useful proteins or metabolic chemicals in plants or to improve other agriculturally relevant characteristics.

Conversely, blocked or reduced expression of a transcription factor may reduce biosynthesis of unwanted compounds or remove an undesirable trait. Therefore, manipulating transcription factor levels in a plant offers tremendous potential in agricultural biotechnology for modifying a plant's traits.

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The present invention provides novel transcription factors useful for modifying a plant's phenotype in desirable ways, such as modifying the characteristics of a plant's seed.

SUMMARY OF THE INVENTION

In a first aspect, the invention relates to a recombinant polynucleotide comprising a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence encoding a polypeptide comprising a sequence selected from SEQ ID Nos. 2N, where N=1-27, or a complementary nucleotide sequence thereof; (b) a nucleotide sequence encoding a polypeptide comprising a conservatively substituted variant of a polypeptide of (a); (c) a nucleotide sequence comprising a sequence selected from those of SEQ ID Nos. 2N-1, where N=1-27, or a

complementary nucleotide sequence thereof; (d) a nucleotide sequence comprising silent substitutions in a nucleotide sequence of (c); (e) a nucleotide sequence which hybridizes under stringent conditions over substantially the entire length of a nucleotide sequence of one or more of: (a), (b), (c), or (d); (f) a nucleotide sequence comprising at least 15 consecutive nucleotides of a sequence of any of (a)-(e); (g) a nucleotide sequence comprising a subsequence or fragment of any of (a)-(f), which subsequence or fragment encodes a polypeptide having a biological activity that modifies a plant's seed characteristics; (h) a nucleotide sequence having at least 31% sequence identity to a nucleotide sequence of any of (a)-(g); (i) a nucleotide sequence having at least 60% identity sequence identity to a nucleotide sequence of any of (a)-(g); (j) a nucleotide sequence which encodes a polypeptide having at least 31% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-27; (k) a nucleotide sequence which encodes a polypeptide having at least 60% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-27; and (l) a nucleotide sequence which encodes a conserved domain of a polypeptide having at least 65% sequence identity to a conserved domain of a polypeptide of SEQ ID Nos. 2N, where N=1-27. The recombinant polynucleotide may further comprise a constitutive, inducible, or tissue-active promoter operably linked to the nucleotide sequence. The invention also relates to compositions comprising at least two of the above described polynucleotides.

In a second aspect, the invention is an isolated or recombinant polypeptide comprising a subsequence of at least about 15 contiguous amino acids encoded by the recombinant or isolated polynucleotide described above.

In another aspect, the invention is a transgenic plant comprising one or more of the above described recombinant polynucleotides. In yet another aspect, the invention is a plant with altered expression levels of a polynucleotide described above or a plant with altered expression or activity levels of an above described polypeptide. Further, the invention may be a plant lacking a nucleotide sequence encoding a polypeptide described above. The plant may be a soybean, wheat, corn, potato, cotton, rice, oilseed rape, sunflower, alfalfa, sugarcane, turf, banana, blackberry, blueberry, strawberry, raspberry, cantaloupe, carrot, cauliflower, coffee, cucumber, eggplant, grapes, honeydew, lettuce, mango, melon, onion, papaya, peas, peppers, pineapple, spinach, squash, sweet corn, tobacco, tomato, watermelon, rosaceous fruits, or vegetable brassicas plant.

In a further aspect, the invention relates to a cloning or expression vector comprising the isolated or recombinant polynucleotide described above or cells comprising the cloning or expression vector.

In yet a further aspect, the invention relates to a composition produced by incubating a polynucleotide of the invention with a nuclease, a restriction enzyme, a polymerase; a polymerase and a primer; a cloning vector, or with a cell.

Furthermore, the invention relates to a method for producing a plant having improved seed traits. The method comprises altering the expression of an isolated or recombinant polynucleotide of the invention or altering the expression or activity of a polypeptide of the invention in a plant to produce a modified plant, and selecting the modified plant for modified seed traits.

In another aspect, the invention relates to a method of identifying a factor that is modulated by or interacts with a polypeptide encoded by a polynucleotide of the invention. The method comprises expressing a polypeptide encoded by the polynucleotide in a plant; and identifying at least one factor that is modulated by or interacts with the polypeptide. In one embodiment the method for identifying modulating or interacting factors is by detecting binding by the polypeptide to a promoter sequence, or by detecting interactions between an additional protein and the polypeptide in a yeast two hybrid system, or by detecting expression of a factor by hybridization to a microarray, subtractive hybridization or differential display.

In yet another aspect, the invention is a method of identifying a molecule that modulates activity or expression of a polynucleotide or polypeptide of interest. The method comprises placing the molecule in contact with a plant comprising the polynucleotide or polypeptide encoded by the polynucleotide of the invention and monitoring one or more of the expression level of the polynucleotide in the plant, the expression level of the polypeptide in the plant, and modulation of an activity of the polypeptide in the plant.

In yet another aspect, the invention relates to an integrated system, computer or computer readable medium comprising one or more character strings corresponding to a polynucleotide of the invention, or to a polypeptide encoded by the polynucleotide. The integrated system, computer or computer readable medium may comprise a link between one or more sequence strings to a modified plant seed trait.

In yet another aspect, the invention is a method for identifying a sequence similar or homologous to one or more polynucleotides of the invention, or one or more polypeptides encoded by the polynucleotides. The method comprises providing a sequence database; and, querying the sequence database with one or more target sequences corresponding to the one or more polynucleotides or to the one or more polypeptides to identify one or more sequence members of the database that display sequence similarity or homology to one or more of the one or more target sequences.

The method may further comprise of linking the one or more of the polynucleotides of the invention, or encoded polypeptides, to a modified plant seed characteristics phenotype.

BRIEF DESCRIPTION OF THE DRAWINGS

5 Figure 1 provides a table of exemplary polynucleotide and polypeptide sequences of the invention. The table includes from left to right for each sequence: the SEQ ID No., the internal code reference number (GID), whether the sequence is a polynucleotide or polypeptide sequence, and identification of any conserved domains for the polypeptide sequences.

10 Figure 2 provides a table of exemplary sequences that are homologous to other sequences provided in the Sequence Listing and that are derived from *Arabidopsis thaliana*. The table includes from left to right: the SEQ ID No., the internal code reference number (GID), identification of the homologous sequence, whether the sequence is a polynucleotide or polypeptide sequence, and identification of any conserved domains for the polypeptide sequences.

15 Figure 3 provides a table of exemplary sequences that are homologous to the sequences provided in Figures 1 and 2 and that are derived from plants other than *Arabidopsis thaliana*. The table includes from left to right: the SEQ ID No., the internal code reference number (GID), the unique GenBank sequence ID No. (NID), the probability that the comparison was generated by chance (P-value), and the species from which the homologous gene was identified.

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DETAILED DESCRIPTION

The present invention relates to polynucleotides and polypeptides, e.g. for modifying phenotypes of plants.

In particular, the polynucleotides or polypeptides are useful for modifying traits associated with a plant's seed characteristics when the expression levels of the polynucleotides or expression levels or activity levels of the polypeptides are altered. Specifically, the polynucleotides and polypeptides are useful for modifying the nutritional content or composition of seeds: such as to modify the protein or oil content of seeds, to modify insoluble sugar content or composition, such as by altering the levels of arabinose, fucose, galactose, mannose, rhamnose or xylose or the like; modify prenyl lipid content or composition, such as by altering the levels of lutein, beta-carotene, xanthophyll-1, xanthophyll-2, chlorophylls A or B, or alpha-, delta- or gamma-tocopherol or the like; modify fatty acid content or composition, such as by altering the levels of the fatty acids 16:0 (palmitic acid), 16:1 (palmitoleic acid), 18:0 (stearic acid), 18:1

(oleic acid), 18:2 (linoleic acid), 20:0 , 18:3 (linolenic acid), 20:1 (eicosenoic acid), 20:2 and 22:1 (erucic acid); modify wax composition or content, such as by altering the levels of C29, C31, or C33 alkanes; modify sterol composition or content, such as by altering the levels of brassicasterol, campesterol, stigmasterol, sitosterol or stigmastanol or the like, or modify glucosinolate composition or content.

Other seed characteristics that may be modified include traits relating to a seed's germination characteristics; shelf-life; drydown characteristics; size; stress responses, such as to heat, cold, salt or osmotic shock; other nutritional content, such as vitamins, minerals, or flavonoids; seedling vigor; pest resistance, or seed coat quality, resistance to pathogens, germination rate, resistance to heavy metals and toxins. Yet another desirable phenotype is a change in the overall gene expression pattern of the seed.

The polynucleotides of the invention encode plant transcription factors. The plant transcription factors are derived, e.g., from *Arabidopsis thaliana* and can belong, e.g., to one or more of the following transcription factor families: the AP2 (APETALA2) domain transcription factor family (Riechmann and Meyerowitz (1998) *J. Biol. Chem.* 379:633-646); the MYB transcription factor family (Martin and Paz-Ares (1997) *Trends Genet.* 13:67-73); the MADS domain transcription factor family (Riechmann and Meyerowitz (1997) *J. Biol. Chem.* 378:1079-1101); the WRKY protein family (Ishiguro and Nakamura (1994) *Mol. Gen. Genet.* 244:563-571); the ankyrin-repeat protein family (Zhang et al. (1992) *Plant Cell* 4:1575-1588); the miscellaneous protein (MISC) family (Kim et al. (1997) *Plant J.* 11:1237-1251); the zinc finger protein (Z) family (Klug and Schwabe (1995) *FASEB J.* 9: 597-604); the homeobox (HB) protein family (Duboule (1994) *Guidebook to the Homeobox Genes*, Oxford University Press); the CAAT-element binding proteins (Forsburg and Guarente (1989) *Genes Dev.* 3:1166-1178); the squamosa promoter binding proteins (SPB) (Klein et al. (1996) *Mol. Gen. Genet.* 1996 250:7-16); the NAM protein family; the IAA/AUX proteins (Rouse et al. (1998) *Science* 279:1371-1373); the HLH/MYC protein family (Littlewood et al. (1994) *Prot. Profile* 1:639-709); the DNA-binding protein (DBP) family (Tucker et al. (1994) *EMBO J.* 13:2994-3002); the bZIP family of transcription factors (Foster et al. (1994) *FASEB J.* 8:192-200); the BPF-1 protein (Box P-binding factor) family (da Costa e Silva et al. (1993) *Plant J.* 4:125-135); and the golden protein (GLD) family (Hall et al. (1998) *Plant Cell* 10:925-936).

In addition to methods for modifying a plant phenotype by employing one or more polynucleotides and polypeptides of the invention described herein, the polynucleotides and polypeptides of the invention have a variety of additional uses. These uses include their use in the recombinant production (i.e, expression) of proteins; as regulators of plant gene expression,

as diagnostic probes for the presence of complementary or partially complementary nucleic acids (including for detection of natural coding nucleic acids); as substrates for further reactions, e.g., mutation reactions, PCR reactions, or the like, or as substrates for cloning e.g., including digestion or ligation reactions, and for identifying exogenous or endogenous modulators of the transcription factors.

DEFINITIONS

A "polynucleotide" is a nucleic acid sequence comprising a plurality of polymerized nucleotide residues, e.g., at least about 15 consecutive polymerized nucleotide residues, optionally at least about 30 consecutive nucleotides, at least about 50 consecutive 10 nucleotides. In many instances, a polynucleotide comprises a nucleotide sequence encoding a polypeptide (or protein) or a domain or fragment thereof. Additionally, the polynucleotide may comprise a promoter, an intron, an enhancer region, a polyadenylation site, a translation initiation site, 5' or 3' untranslated regions, a reporter gene, a selectable marker, or the like. The polynucleotide can be single stranded or double stranded DNA or RNA. The polynucleotide 15 optionally comprises modified bases or a modified backbone. The polynucleotide can be, e.g., genomic DNA or RNA, a transcript (such as an mRNA), a cDNA, a PCR product, a cloned DNA, a synthetic DNA or RNA, or the like. The polynucleotide can comprise a sequence in either sense or antisense orientations.

A "recombinant polynucleotide" is a polynucleotide that is not in its native state, 20 e.g., the polynucleotide comprises a nucleotide sequence not found in nature, or the polynucleotide is in a context other than that in which it is naturally found, e.g., separated from nucleotide sequences with which it typically is in proximity in nature, or adjacent (or contiguous with) nucleotide sequences with which it typically is not in proximity. For example, the sequence at issue can be cloned into a vector, or otherwise recombined with one or more additional nucleic 25 acid.

An "isolated polynucleotide" is a polynucleotide whether naturally occurring or recombinant, that is present outside the cell in which it is typically found in nature, whether purified or not. Optionally, an isolated polynucleotide is subject to one or more enrichment or purification procedures, e.g., cell lysis, extraction, centrifugation, precipitation, or the like.

30 A "recombinant polypeptide" is a polypeptide produced by translation of a recombinant polynucleotide. An "isolated polypeptide," whether a naturally occurring or a recombinant polypeptide, is more enriched in (or out of) a cell than the polypeptide in its natural state in a wild type cell, e.g., more than about 5% enriched, more than about 10% enriched, or

more than about 20%, or more than about 50%, or more, enriched, i.e., alternatively denoted: 105%, 110%, 120%, 150% or more, enriched relative to wild type standardized at 100%. Such an enrichment is not the result of a natural response of a wild type plant. Alternatively, or 5 additionally, the isolated polypeptide is separated from other cellular components with which it is typically associated, e.g., by any of the various protein purification methods herein.

The term "transgenic plant" refers to a plant that contains genetic material, not found in a wild type plant of the same species, variety or cultivar. The genetic material may include a transgene, an insertional mutagenesis event (such as by transposon or T-DNA insertional mutagenesis), an activation tagging sequence, a mutated sequence, a homologous 10 recombination event or a sequence modified by chimeroplasty. Typically, the foreign genetic material has been introduced into the plant by human manipulation.

A transgenic plant may contain an expression vector or cassette. The expression cassette typically comprises a polypeptide-encoding sequence operably linked (i.e., under regulatory control of) to appropriate inducible or constitutive regulatory sequences that allow for 15 the expression of polypeptide. The expression cassette can be introduced into a plant by transformation or by breeding after transformation of a parent plant. A plant refers to a whole plant as well as to a plant part, such as seed, fruit, leaf, or root, plant tissue, plant cells or any other plant material, e.g., a plant explant, as well as to progeny thereof, and to *in vitro* systems that mimic biochemical or cellular components or processes in a cell.

20 The phrase "ectopically expression or altered expression" in reference to a polynucleotide indicates that the pattern of expression in, e.g., a transgenic plant or plant tissue, is different from the expression pattern in a wild type plant or a reference plant of the same species. For example, the polynucleotide or polypeptide is expressed in a cell or tissue type other than a 25 cell or tissue type in which the sequence is expressed in the wild type plant, or by expression at a time other than at the time the sequence is expressed in the wild type plant, or by a response to different inducible agents, such as hormones or environmental signals, or at different expression levels (either higher or lower) compared with those found in a wild type plant. The term also refers to altered expression patterns that are produced by lowering the levels of expression to below the detection level or completely abolishing expression. The resulting expression pattern 30 can be transient or stable, constitutive or inducible. In reference to a polypeptide, the term "ectopic expression or altered expression" further may relate to altered activity levels resulting from the interactions of the polypeptides with exogenous or endogenous modulators or from interactions with factors or as a result of the chemical modification of the polypeptides.

The term "fragment" or "domain," with respect to a polypeptide, refers to a subsequence of the polypeptide. In some cases, the fragment or domain, is a subsequence of the polypeptide which performs at least one biological function of the intact polypeptide in substantially the same manner, or to a similar extent, as does the intact polypeptide. For example,

5 a polypeptide fragment can comprise a recognizable structural motif or functional domain such as a DNA binding domain that binds to a DNA promoter region, an activation domain or a domain for protein-protein interactions. Fragments can vary in size from as few as 6 amino acids to the full length of the intact polypeptide, but are preferably at least about 30 amino acids in length and more preferably at least about 60 amino acids in length. In reference to a nucleotide sequence, "a

10 fragment" refers to any subsequence of a polynucleotide, typically, of at least consecutive about 15 nucleotides, preferably at least about 30 nucleotides, more preferably at least about 50, of any of the sequences provided herein.

The term "trait" refers to a physiological, morphological, biochemical or physical characteristic of a plant or particular plant material or cell. In some instances, this characteristic

15 is visible to the human eye, such as seed or plant size, or can be measured by available biochemical techniques, such as the protein, starch or oil content of seed or leaves or by the observation of the expression level of genes, e.g., by employing Northern analysis, RT-PCR, microarray gene expression assays or reporter gene expression systems, or by agricultural observations such as stress tolerance, yield or pathogen tolerance.

20 "Trait modification" refers to a detectable difference in a characteristic in a plant ectopically expressing a polynucleotide or polypeptide of the present invention relative to a plant not doing so, such as a wild type plant. In some cases, the trait modification can be evaluated quantitatively. For example, the trait modification can entail at least about a 2% increase or decrease in an observed trait (difference), at least a 5% difference, at least about a 10%

25 difference, at least about a 20% difference, at least about a 30%, at least about a 50%, at least about a 70%, or at least about a 100%, or an even greater difference. It is known that there can be a natural variation in the modified trait. Therefore, the trait modification observed entails a change of the normal distribution of the trait in the plants compared with the distribution observed in wild type plant.

30 Trait modifications of particular interest include those to seed (such as embryo or endosperm), fruit, root, flower, leaf, stem, shoot, seedling or the like, including: enhanced tolerance to environmental conditions including freezing, chilling, heat, drought, water saturation, radiation and ozone; improved tolerance to microbial, fungal or viral diseases; improved tolerance to pest infestations, including nematodes, mollicutes, parasitic higher plants or the like;

decreased herbicide sensitivity; improved tolerance of heavy metals or enhanced ability to take up heavy metals; improved growth under poor photoconditions (e.g., low light and/or short day length), or changes in expression levels of genes of interest. Other phenotype that can be modified relate to the production of plant metabolites, such as variations in the production of 5 taxol, tocopherol, tocotrienol, sterols, phytosterols, vitamins, wax monomers, anti-oxidants, amino acids, lignins, cellulose, tannins, prenyllipids (such as chlorophylls and carotenoids), glucosinolates, and terpenoids, enhanced or compositionally altered protein or oil production (especially in seeds), or modified sugar (insoluble or soluble) and/or starch composition.

Physical plant characteristics that can be modified include cell development (such as the number 10 of trichomes), fruit and seed size and number, yields of plant parts such as stems, leaves and roots, the stability of the seeds during storage, characteristics of the seed pod (e.g., susceptibility to shattering), root hair length and quantity, internode distances, or the quality of seed coat. Plant growth characteristics that can be modified include growth rate, germination rate of seeds, vigor 15 of plants and seedlings, leaf and flower senescence, male sterility, apomixis, flowering time, flower abscission, rate of nitrogen uptake, biomass or transpiration characteristics, as well as plant architecture characteristics such as apical dominance, branching patterns, number of organs, organ identity, organ shape or size.

POLYPEPTIDES AND POLYNUCLEOTIDES OF THE INVENTION

The present invention provides, among other things, transcription factors (TFs), 20 and transcription factor homologue polypeptides, and isolated or recombinant polynucleotides encoding the polypeptides. These polypeptides and polynucleotides may be employed to modify a plant's seed characteristics.

Exemplary polynucleotides encoding the polypeptides of the invention were identified in the *Arabidopsis thaliana* GenBank database using publicly available sequence 25 analysis programs and parameters. Sequences initially identified were then further characterized to identify sequences comprising specified sequence strings corresponding to sequence motifs present in families of known transcription factors. Polynucleotide sequences meeting such criteria were confirmed as transcription factors.

Additional polynucleotides of the invention were identified by screening 30 *Arabidopsis thaliana* and/or other plant cDNA libraries with probes corresponding to known transcription factors under low stringency hybridization conditions. Additional sequences, including full length coding sequences were subsequently recovered by the rapid amplification of cDNA ends (RACE) procedure, using a commercially available kit according to the

manufacturer's instructions. Where necessary, multiple rounds of RACE are performed to isolate 5' and 3' ends. The full length cDNA was then recovered by a routine end-to-end polymerase chain reaction (PCR) using primers specific to the isolated 5' and 3' ends. Exemplary sequences are provided in the Sequence Listing.

5 The polynucleotides of the invention were ectopically expressed in overexpressor or knockout plants and changes in the seed characteristics of the plants were observed. Therefore, the polynucleotides and polypeptides can be employed to improve the seed characteristics of plants.

10 Making polynucleotides
10 The polynucleotides of the invention include sequences that encode transcription factors and transcription factor homologue polypeptides and sequences complementary thereto, as well as unique fragments of coding sequence, or sequence complementary thereto. Such polynucleotides can be, e.g., DNA or RNA, e.g., mRNA, cRNA, synthetic RNA, genomic DNA, cDNA synthetic DNA, oligonucleotides, etc. The polynucleotides are either double-stranded or
15 single-stranded, and include either, or both sense (i.e., coding) sequences and antisense (i.e., non-coding, complementary) sequences. The polynucleotides include the coding sequence of a transcription factor, or transcription factor homologue polypeptide, in isolation, in combination with additional coding sequences (e.g., a purification tag, a localization signal, as a fusion-protein, as a pre-protein, or the like), in combination with non-coding sequences (e.g., introns or
20 inteins, regulatory elements such as promoters, enhancers, terminators, and the like), and/or in a vector or host environment in which the polynucleotide encoding a transcription factor or transcription factor homologue polypeptide is an endogenous or exogenous gene.

25 A variety of methods exist for producing the polynucleotides of the invention. Procedures for identifying and isolating DNA clones are well known to those of skill in the art, and are described in, e.g., Berger and Kimmel, Guide to Molecular Cloning Techniques, Methods in Enzymology volume 152 Academic Press, Inc., San Diego, CA ("Berger"); Sambrook et al., Molecular Cloning - A Laboratory Manual (2nd Ed.), Vol. 1-3, Cold Spring Harbor Laboratory, Cold Spring Harbor, New York, 1989 ("Sambrook") and Current Protocols in Molecular Biology, F.M. Ausubel et al., eds., Current Protocols, a joint venture between Greene Publishing
30 Associates, Inc. and John Wiley & Sons, Inc., (supplemented through 2000) ("Ausubel").

35 Alternatively, polynucleotides of the invention, can be produced by a variety of in vitro amplification methods adapted to the present invention by appropriate selection of specific or degenerate primers. Examples of protocols sufficient to direct persons of skill through in vitro amplification methods, including the polymerase chain reaction (PCR) the ligase chain

reaction (LCR), Qbeta-replicase amplification and other RNA polymerase mediated techniques (e.g., NASBA), e.g., for the production of the homologous nucleic acids of the invention are found in Berger, Sambrook, and Ausubel, as well as Mullis et al., (1987) PCR Protocols A Guide to Methods and Applications (Innis et al. eds) Academic Press Inc. San Diego, CA (1990) (Innis).
5 Improved methods for cloning in vitro amplified nucleic acids are described in Wallace et al., U.S. Pat. No. 5,426,039. Improved methods for amplifying large nucleic acids by PCR are summarized in Cheng et al. (1994) Nature 369: 684-685 and the references cited therein, in which PCR amplicons of up to 40kb are generated. One of skill will appreciate that essentially any RNA can be converted into a double stranded DNA suitable for restriction digestion, PCR
10 expansion and sequencing using reverse transcriptase and a polymerase. See, e.g., Ausubel, Sambrook and Berger, *all supra*.

Alternatively, polynucleotides and oligonucleotides of the invention can be assembled from fragments produced by solid-phase synthesis methods. Typically, fragments of up to approximately 100 bases are individually synthesized and then enzymatically or chemically
15 ligated to produce a desired sequence, e.g., a polynucleotide encoding all or part of a transcription factor. For example, chemical synthesis using the phosphoramidite method is described, e.g., by Beaucage et al. (1981) Tetrahedron Letters 22:1859-69; and Matthes et al. (1984) EMBO J. 3:801-5. According to such methods, oligonucleotides are synthesized, purified, annealed to their complementary strand, ligated and then optionally cloned into suitable vectors.
20 And if so desired, the polynucleotides and polypeptides of the invention can be custom ordered from any of a number of commercial suppliers.

HOMOLOGOUS SEQUENCES

Sequences homologous, i.e., that share significant sequence identity or similarity, to those provided in the Sequence Listing, derived from *Arabidopsis thaliana* or from other plants
25 of choice are also an aspect of the invention. Homologous sequences can be derived from any plant including monocots and dicots and in particular agriculturally important plant species, including but not limited to, crops such as soybean, wheat, corn, potato, cotton, rice, oilseed rape (including canola), sunflower, alfalfa, sugarcane and turf; or fruits and vegetables, such as banana, blackberry, blueberry, strawberry, and raspberry, cantaloupe, carrot, cauliflower, coffee,
30 cucumber, eggplant, grapes, honeydew, lettuce, mango, melon, onion, papaya, peas, peppers, pineapple, spinach, squash, sweet corn, tobacco, tomato, watermelon, rosaceous fruits (such as apple, peach, pear, cherry and plum) and vegetable brassicas (such as broccoli, cabbage, cauliflower, brussel sprouts and kohlrabi). Other crops, fruits and vegetables whose phenotype

can be changed include barley, rye, millet, sorghum, currant, avocado, citrus fruits such as oranges, lemons, grapefruit and tangerines, artichoke, cherries, nuts such as the walnut and peanut, endive, leek, roots, such as arrowroot, beet, cassava, turnip, radish, yam, and sweet potato, and beans. The homologous sequences may also be derived from woody species, such 5 pine, poplar and eucalyptus.

- Transcription factors that are homologous to the listed sequences will typically share at least about 31% amino acid sequence identity. More closely related transcription factors can share at least about 50%, about 60%, about 65%, about 70%, about 75% or about 80% or about 90% or about 95% or about 98% or more sequence identity with the listed sequences.
- 10 Factors that are most closely related to the listed sequences share, e.g., at least about 85%, about 90% or about 95% or more % sequence identity to the listed sequences. At the nucleotide level, the sequences will typically share at least about 40% nucleotide sequence identity, preferably at least about 50%, about 60%, about 70% or about 80% sequence identity, and more preferably about 85%, about 90%, about 95% or about 97% or more sequence identity to one or more of the 15 listed sequences. The degeneracy of the genetic code enables major variations in the nucleotide sequence of a polynucleotide while maintaining the amino acid sequence of the encoded protein. Conserved domains within a transcription factor family may exhibit a higher degree of sequence homology, such as at least 65% sequence identity including conservative substitutions, and preferably at least 80% sequence identity.

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Identifying Nucleic Acids by Hybridization

Polynucleotides homologous to the sequences illustrated in the Sequence Listing can be identified, e.g., by hybridization to each other under stringent or under highly stringent conditions. Single stranded polynucleotides hybridize when they associate based on a variety of well characterized physico-chemical forces, such as hydrogen bonding, solvent exclusion, base 25 stacking and the like. The stringency of a hybridization reflects the degree of sequence identity of the nucleic acids involved, such that the higher the stringency, the more similar are the two polynucleotide strands. Stringency is influenced by a variety of factors, including temperature, salt concentration and composition, organic and non-organic additives, solvents, etc. present in both the hybridization and wash solutions and incubations (and number), as described in more 30 detail in the references cited above.

An example of stringent hybridization conditions for hybridization of complementary nucleic acids which have more than 100 complementary residues on a filter in a Southern or northern blot is about 5°C to 20°C lower than the thermal melting point (T_m) for the specific sequence at a defined ionic strength and pH. The T_m is the temperature (under defined

ionic strength and pH) at which 50% of the target sequence hybridizes to a perfectly matched probe. Nucleic acid molecules that hybridize under stringent conditions will typically hybridize to a probe based on either the entire cDNA or selected portions, e.g., to a unique subsequence, of the cDNA under wash conditions of 0.2x SSC to 2.0 x SSC, 0.1% SDS at 50-65° C, for example 5 0.2 x SSC, 0.1% SDS at 65° C. For identification of less closely related homologues washes can be performed at a lower temperature, e.g., 50° C. In general, stringency is increased by raising the wash temperature and/or decreasing the concentration of SSC.

As another example, stringent conditions can be selected such that an oligonucleotide that is perfectly complementary to the coding oligonucleotide hybridizes to the 10 coding oligonucleotide with at least about a 5-10x higher signal to noise ratio than the ratio for hybridization of the perfectly complementary oligonucleotide to a nucleic acid encoding a transcription factor known as of the filing date of the application. Conditions can be selected such that a higher signal to noise ratio is observed in the particular assay which is used, e.g., about 15x, 25x, 35x, 50x or more. Accordingly, the subject nucleic acid hybridizes to the unique 15 coding oligonucleotide with at least a 2x higher signal to noise ratio as compared to hybridization of the coding oligonucleotide to a nucleic acid encoding known polypeptide. Again, higher signal to noise ratios can be selected, e.g., about 5x, 10x, 25x, 35x, 50x or more. The particular signal will depend on the label used in the relevant assay, e.g., a fluorescent label, a colorimetric label, a radio active label, or the like.

20 Alternatively, transcription factor homologue polypeptides can be obtained by screening an expression library using antibodies specific for one or more transcription factors. With the provision herein of the disclosed transcription factor, and transcription factor homologue nucleic acid sequences, the encoded polypeptide(s) can be expressed and purified in a heterologous expression system (e.g., *E. coli*) and used to raise antibodies (monoclonal or 25 polyclonal) specific for the polypeptide(s) in question. Antibodies can also be raised against synthetic peptides derived from transcription factor, or transcription factor homologue, amino acid sequences. Methods of raising antibodies are well known in the art and are described in Harlow and Lane (1988) Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, New York. Such antibodies can then be used to screen an expression library produced from the plant 30 from which it is desired to clone additional transcription factor homologues, using the methods described above. The selected cDNAs can be confirmed by sequencing and enzymatic activity.

SEQUENCE VARIATIONS

It will readily be appreciated by those of skill in the art, that any of a variety of polynucleotide sequences are capable of encoding the transcription factors and transcription factor homologue polypeptides of the invention. Due to the degeneracy of the genetic code, 5 many different polynucleotides can encode identical and/or substantially similar polypeptides in addition to those sequences illustrated in the Sequence Listing.

For example, Table 1 illustrates, e.g., that the codons AGC, AGT, TCA, TCC, TCG, and TCT all encode the same amino acid: serine. Accordingly, at each position in the sequence where there is a codon encoding serine, any of the above trinucleotide sequences can be 10 used without altering the encoded polypeptide.

Table 1

Amino acids			Codon					
Alanine	Ala	A	GCA	GCC	GCG	GCU		
Cysteine	Cys	C	TGC	TGT				
Aspartic acid	Asp	D	GAC	GAT				
Glutamic acid	Glu	E	GAA	GAG				
Phenylalanine	Phe	F	TTC	TTT				
Glycine	Gly	G	GGA	GGC	GGG	GGT		
Histidine	His	H	CAC	CAT				
Isoleucine	Ile	I	ATA	ATC	ATT			
Lysine	Lys	K	AAA	AAG				
Leucine	Leu	L	TTA	TTG	CTA	CTC	CTG	CTT
Methionine	Met	M	ATG					
Asparagine	Asn	N	AAC	AAT				
Proline	Pro	P	CCA	CCC	CCG	CCT		
Glutamine	Gln	Q	CAA	CAG				
Arginine	Arg	R	AGA	AGG	CGA	CGC	CGG	CGT
Serine	Ser	S	AGC	AGT	TCA	TCC	TCG	TCT
Threonine	Thr	T	ACA	ACC	ACG	ACT		
Valine	Val	V	GTA	GTC	GTG	GTT		
Tryptophan	Trp	W	TGG					
Tyrosine	Tyr	Y	TAC	TAT				

Sequence alterations that do not change the amino acid sequence encoded by the 15 polynucleotide are termed "silent" variations. With the exception of the codons ATG and TGG, encoding methionine and tryptophan, respectively, any of the possible codons for the same amino acid can be substituted by a variety of techniques, e.g., site-directed mutagenesis, available in the art. Accordingly, any and all such variations of a sequence selected from the above table are a feature of the invention.

In addition to silent variations, other conservative variations that alter one, or a few amino acids in the encoded polypeptide, can be made without altering the function of the polypeptide, these conservative variants are, likewise, a feature of the invention.

For example, substitutions, deletions and insertions introduced into the sequences provided in the Sequence Listing are also envisioned by the invention. Such sequence modifications can be engineered into a sequence by site-directed mutagenesis (Wu (ed.) Meth. Enzymol. (1993) vol. 217, Academic Press) or the other methods noted below. Amino acid substitutions are typically of single residues; insertions usually will be on the order of about from 1 to 10 amino acid residues; and deletions will range about from 1 to 30 residues. In preferred embodiments, deletions or insertions are made in adjacent pairs, e.g., a deletion of two residues or insertion of two residues. Substitutions, deletions, insertions or any combination thereof can be combined to arrive at a sequence. The mutations that are made in the polynucleotide encoding the transcription factor should not place the sequence out of reading frame and should not create complementary regions that could produce secondary mRNA structure. Preferably, the polypeptide encoded by the DNA performs the desired function.

Conservative substitutions are those in which at least one residue in the amino acid sequence has been removed and a different residue inserted in its place. Such substitutions generally are made in accordance with the Table 2 when it is desired to maintain the activity of the protein. Table 2 shows amino acids which can be substituted for an amino acid in a protein and which are typically regarded as conservative substitutions.

25

30

Table 2

Residue	Conservative Substitutions
Ala	Ser
Arg	Lys
Asn	Gln; His
Asp	Glu
Gln	Asn
Cys	Ser
Glu	Asp
Gly	Pro
His	Asn; Gln
Ile	Leu, Val
Leu	Ile; Val
Lys	Arg; Gln
Met	Leu; Ile
Phe	Met; Leu; Tyr
Ser	Thr; Gly
Thr	Ser; Val
Trp	Tyr
Tyr	Trp; Phe
Val	Ile; Leu

Substitutions that are less conservative than those in Table 2 can be selected by picking residues that differ more significantly in their effect on maintaining (a) the structure of the polypeptide backbone in the area of the substitution, for example, as a sheet or helical conformation, (b) the charge or hydrophobicity of the molecule at the target site, or (c) the bulk of the side chain. The substitutions which in general are expected to produce the greatest changes in protein properties will be those in which (a) a hydrophilic residue, e.g., seryl or thronyl, is substituted for (or by) a hydrophobic residue, e.g., leucyl, isoleucyl, phenylalanyl, valyl or alanyl; (b) a cysteine or proline is substituted for (or by) any other residue; (c) a residue having an electropositive side chain, e.g., lysyl, arginyl, or histidyl, is substituted for (or by) an electronegative residue, e.g., glutamyl or aspartyl; or (d) a residue having a bulky side chain, e.g., phenylalanine, is substituted for (or by) one not having a side chain, e.g., glycine.

FURTHER MODIFYING SEQUENCES OF THE INVENTION—MUTATION/ FORCED
EVOLUTION

In addition to generating silent or conservative substitutions as noted, above, the present invention optionally includes methods of modifying the sequences of the Sequence Listing.

- 5 Listing. In the methods, nucleic acid or protein modification methods are used to alter the given sequences to produce new sequences and/or to chemically or enzymatically modify given sequences to change the properties of the nucleic acids or proteins.

Thus, in one embodiment, given nucleic acid sequences are modified, e.g., according to standard mutagenesis or artificial evolution methods to produce modified sequences.

- 10 For example, Ausubel, *supra*, provides additional details on mutagenesis methods. Artificial forced evolution methods are described, e.g., by Stemmer (1994) *Nature* 370:389-391, and Stemmer (1994) *Proc. Natl. Acad. Sci. USA* 91:10747-10751. Many other mutation and evolution methods are also available and expected to be within the skill of the practitioner.

- 15 Similarly, chemical or enzymatic alteration of expressed nucleic acids and polypeptides can be performed by standard methods. For example, sequence can be modified by addition of lipids, sugars, peptides, organic or inorganic compounds, by the inclusion of modified nucleotides or amino acids, or the like. For example, protein modification techniques are illustrated in Ausubel, *supra*. Further details on chemical and enzymatic modifications can be found herein. These modification methods can be used to modify any given sequence, or to 20 modify any sequence produced by the various mutation and artificial evolution modification methods noted herein.

- Accordingly, the invention provides for modification of any given nucleic acid by mutation, evolution, chemical or enzymatic modification, or other available methods, as well as for the products produced by practicing such methods, e.g., using the sequences herein as a 25 starting substrate for the various modification approaches.

- For example, optimized coding sequence containing codons preferred by a particular prokaryotic or eukaryotic host can be used e.g., to increase the rate of translation or to produce recombinant RNA transcripts having desirable properties, such as a longer half-life, as compared with transcripts produced using a non-optimized sequence. Translation stop codons 30 can also be modified to reflect host preference. For example, preferred stop codons for *S. cerevisiae* and mammals are TAA and TGA, respectively. The preferred stop codon for monocotyledonous plants is TGA, whereas insects and *E. coli* prefer to use TAA as the stop codon.

The polynucleotide sequences of the present invention can also be engineered in order to alter a coding sequence for a variety of reasons, including but not limited to, alterations which modify the sequence to facilitate cloning, processing and/or expression of the gene product. For example, alterations are optionally introduced using techniques which are well known in the art, e.g., site-directed mutagenesis, to insert new restriction sites, to alter glycosylation patterns, to change codon preference, to introduce splice sites, etc.

Furthermore, a fragment or domain derived from any of the polypeptides of the invention can be combined with domains derived from other transcription factors or synthetic domains to modify the biological activity of a transcription factor. For instance, a DNA binding domain derived from a transcription factor of the invention can be combined with the activation domain of another transcription factor or with a synthetic activation domain. A transcription activation domain assists in initiating transcription from a DNA binding site. Examples include the transcription activation region of VP16 or GAL4 (Moore et al. (1998) Proc. Natl. Acad. Sci. USA 95: 376-381; and Aoyama et al. (1995) Plant Cell 7:1773-1785), peptides derived from bacterial sequences (Ma and Ptashne (1987) Cell 51: 113-119) and synthetic peptides (Giniger and Ptashne, (1987) Nature 330:670-672).

EXPRESSION AND MODIFICATION OF POLYPEPTIDES

Typically, polynucleotide sequences of the invention are incorporated into recombinant DNA (or RNA) molecules that direct expression of polypeptides of the invention in appropriate host cells, transgenic plants, in vitro translation systems, or the like. Due to the inherent degeneracy of the genetic code, nucleic acid sequences which encode substantially the same or a functionally equivalent amino acid sequence can be substituted for any listed sequence to provide for cloning and expressing the relevant homologue.

Vectors, Promoters and Expression Systems

The present invention includes recombinant constructs comprising one or more of the nucleic acid sequences herein. The constructs typically comprise a vector, such as a plasmid, a cosmid, a phage, a virus (e.g., a plant virus), a bacterial artificial chromosome (BAC), a yeast artificial chromosome (YAC), or the like, into which a nucleic acid sequence of the invention has been inserted, in a forward or reverse orientation. In a preferred aspect of this embodiment, the construct further comprises regulatory sequences, including, for example, a promoter, operably linked to the sequence. Large numbers of suitable vectors and promoters are known to those of skill in the art, and are commercially available.

General texts which describe molecular biological techniques useful herein, including the use and production of vectors, promoters and many other relevant topics, include Berger, Sambrook and Ausubel, *supra*. Any of the identified sequences can be incorporated into a cassette or vector, e.g., for expression in plants. A number of expression vectors suitable for stable transformation of plant cells or for the establishment of transgenic plants have been described including those described in Weissbach and Weissbach, (1989) Methods for Plant Molecular Biology, Academic Press, and Gelvin et al., (1990) Plant Molecular Biology Manual, Kluwer Academic Publishers. Specific examples include those derived from a Ti plasmid of *Agrobacterium tumefaciens*, as well as those disclosed by Herrera-Estrella et al. (1983) Nature 303: 209, Bevan (1984) Nucl Acid Res. 12: 8711-8721, Klee (1985) Bio/Technology 3: 637-642, for dicotyledonous plants.

Alternatively, non-Ti vectors can be used to transfer the DNA into monocotyledonous plants and cells by using free DNA delivery techniques. Such methods can involve, for example, the use of liposomes, electroporation, microprojectile bombardment, silicon carbide whiskers, and viruses. By using these methods transgenic plants such as wheat, rice (Christou (1991) Bio/Technology 9: 957-962) and corn (Gordon-Kamm (1990) Plant Cell 2: 603-618) can be produced. An immature embryo can also be a good target tissue for monocots for direct DNA delivery techniques by using the particle gun. (Weeks et al. (1993) Plant Physiol 102: 1077-1084; Vasil (1993) Bio/Technology 10: 667-674; Wan and Lemeaux (1994) Plant Physiol 104: 37-48, and for Agrobacterium-mediated DNA transfer (Ishida et al. (1996) Nature Biotech 14: 745-750).

Typically, plant transformation vectors include one or more cloned plant coding sequence (genomic or cDNA) under the transcriptional control of 5' and 3' regulatory sequences and a dominant selectable marker. Such plant transformation vectors typically also contain a promoter (e.g., a regulatory region controlling inducible or constitutive, environmentally-or developmentally-regulated, or cell- or tissue-specific expression), a transcription initiation start site, an RNA processing signal (such as intron splice sites), a transcription termination site, and/or a polyadenylation signal.

Examples of constitutive plant promoters which can be useful for expressing the TF sequence include: the cauliflower mosaic virus (CaMV) 35S promoter, which confers constitutive, high-level expression in most plant tissues (*see, e.g.*, Odel et al. (1985) Nature 313:810); the nopaline synthase promoter (An et al. (1988) Plant Physiol 88:547); and the octopine synthase promoter (Fromm et al. (1989) Plant Cell 1: 977).

A variety of plant gene promoters that regulate gene expression in response to environmental, hormonal, chemical, developmental signals, and in a tissue-active manner can be used for expression of a TF sequence in plants. Choice of a promoter is based largely on the phenotype of interest and is determined by such factors as tissue (e.g., seed, fruit, root, pollen, 5 vascular tissue, flower, carpel, etc.), inducibility (e.g., in response to wounding, heat, cold, drought, light, pathogens, etc.), timing, developmental stage, and the like. Numerous known promoters have been characterized and can favorable be employed to promote expression of a polynucleotide of the invention in a transgenic plant or cell of interest. For example, tissue specific promoters include: seed-specific promoters (such as the napin, phaseolin or DC3 10 promoter described in US Pat. No. 5,773,697), fruit-specific promoters that are active during fruit ripening (such as the dru 1 promoter (US Pat. No. 5,783,393), or the 2A11 promoter (US Pat. No. 4,943,674) and the tomato polygalacturonase promoter (Bird et al. (1988) Plant Mol Biol 11:651), root-specific promoters, such as those disclosed in US Patent Nos. 5,618,988, 5,837,848 and 5,905,186, pollen-active promoters such as PTA29, PTA26 and PTA13 (US Pat. No. 5,792,929), 15 promoters active in vascular tissue (Ringli and Keller (1998) Plant Mol Biol 37:977-988), flower-specific (Kaiser et al, (1995) Plant Mol Biol 28:231-243), pollen (Baerson et al. (1994) Plant Mol Biol 26:1947-1959), carpels (Ohl et al. (1990) Plant Cell 2:837-848), pollen and ovules (Baerson et al. (1993) Plant Mol Biol 22:255-267), auxin-inducible promoters (such as that described in van der Kop et al. (1999) Plant Mol Biol 39:979-990 or Baumann et al. (1999) Plant Cell 11:323- 20 334), cytokinin-inducible promoter (Guevara-Garcia (1998) Plant Mol Biol 38:743-753), promoters responsive to gibberellin (Shi et al. (1998) Plant Mol Biol 38:1053-1060, Willmott et al. (1998) 38:817-825) and the like. Additional promoters are those that elicit expression in response to heat (Ainley et al. (1993) Plant Mol Biol 22: 13-23), light (e.g., the pea rbcS-3A promoter, Kuhlemeier et al. (1989) Plant Cell 1:471, and the maize rbcS promoter, Schaffner and 25 Sheen (1991) Plant Cell 3: 997); wounding (e.g., *wunf*, Siebertz et al. (1989) Plant Cell 1: 961); pathogens (such as the PR-1 promoter described in Buchel et al. (1999) Plant Mol. Biol. 40:387-396, and the PDF1.2 promoter described in Manners et al. (1998) Plant Mol. Biol. 38:1071-80), and chemicals such as methyl jasmonate or salicylic acid (Gatz et al. (1997) Plant Mol Biol 48: 89-108). In addition, the timing of the expression can be controlled by using promoters such as those 30 acting at senescence (An and Amazon (1995) Science 270: 1986-1988); or late seed development (Odell et al. (1994) Plant Physiol 106:447-458).

Plant expression vectors can also include RNA processing signals that can be positioned within, upstream or downstream of the coding sequence. In addition, the expression vectors can include additional regulatory sequences from the 3'-untranslated region of plant

genes, e.g., a 3' terminator region to increase mRNA stability of the mRNA, such as the PI-II terminator region of potato or the octopine or nopaline synthase 3' terminator regions.

Additional Expression Elements

Specific initiation signals can aid in efficient translation of coding sequences.

- 5 These signals can include, e.g., the ATG initiation codon and adjacent sequences. In cases where a coding sequence, its initiation codon and upstream sequences are inserted into the appropriate expression vector, no additional translational control signals may be needed. However, in cases where only coding sequence (e.g., a mature protein coding sequence), or a portion thereof, is inserted, exogenous transcriptional control signals including the ATG initiation codon can be
- 10 separately provided. The initiation codon is provided in the correct reading frame to facilitate transcription. Exogenous transcriptional elements and initiation codons can be of various origins, both natural and synthetic. The efficiency of expression can be enhanced by the inclusion of enhancers appropriate to the cell system in use.

Expression Hosts

- 15 The present invention also relates to host cells which are transduced with vectors of the invention, and the production of polypeptides of the invention (including fragments thereof) by recombinant techniques. Host cells are genetically engineered (i.e., nucleic acids are introduced, e.g.; transduced, transformed or transfected) with the vectors of this invention, which may be, for example, a cloning vector or an expression vector comprising the relevant nucleic acids herein. The vector is optionally a plasmid, a viral particle, a phage, a naked nucleic acids, etc. The engineered host cells can be cultured in conventional nutrient media modified as appropriate for activating promoters, selecting transformants, or amplifying the relevant gene. The culture conditions, such as temperature, pH and the like, are those previously used with the host cell selected for expression, and will be apparent to those skilled in the art and in the 20 references cited herein, including, Sambrook and Ausubel.

- 25 The host cell can be a eukaryotic cell, such as a yeast cell, or a plant cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. Plant protoplasts are also suitable for some applications. For example, the DNA fragments are introduced into plant tissues, cultured plant cells or plant protoplasts by standard methods including electroporation (Fromm et al., 30 (1985) Proc. Natl. Acad. Sci. USA 82, 5824, infection by viral vectors such as cauliflower mosaic virus (CaMV) (Hohn et al., (1982) Molecular Biology of Plant Tumors, (Academic Press, New York) pp. 549-560; US 4,407,956), high velocity ballistic penetration by small particles with the nucleic acid either within the matrix of small beads or particles, or on the surface (Klein et al., (1987) Nature 327, 70-73), use of pollen as vector (WO 85/01856), or use of Agrobacterium

tumefaciens or *A. rhizogenes* carrying a T-DNA plasmid in which DNA fragments are cloned. The T-DNA plasmid is transmitted to plant cells upon infection by *Agrobacterium tumefaciens*, and a portion is stably integrated into the plant genome (Horsch et al. (1984) Science 233:496-498; Fraley et al. (1983) Proc. Natl. Acad. Sci. USA 80, 4803).

5 The cell can include a nucleic acid of the invention which encodes a polypeptide, wherein the cells expresses a polypeptide of the invention. The cell can also include vector sequences, or the like. Furthermore, cells and transgenic plants which include any polypeptide or nucleic acid above or throughout this specification, e.g., produced by transduction of a vector of the invention, are an additional feature of the invention.

10 For long-term, high-yield production of recombinant proteins, stable expression can be used. Host cells transformed with a nucleotide sequence encoding a polypeptide of the invention are optionally cultured under conditions suitable for the expression and recovery of the encoded protein from cell culture. The protein or fragment thereof produced by a recombinant cell may be secreted, membrane-bound, or contained intracellularly, depending on the sequence and/or the vector used. As will be understood by those of skill in the art, expression vectors containing polynucleotides encoding mature proteins of the invention can be designed with signal sequences which direct secretion of the mature polypeptides through a prokaryotic or eukaryotic cell membrane.

20 Modified Amino Acids
Polypeptides of the invention may contain one or more modified amino acids. The presence of modified amino acids may be advantageous in, for example, increasing polypeptide half-life, reducing polypeptide antigenicity or toxicity, increasing polypeptide storage stability, or the like. Amino acid(s) are modified, for example, co-translationally or post-translationally during recombinant production or modified by synthetic or chemical means.

25 Non-limiting examples of a modified amino acid include incorporation or other use of acetylated amino acids, glycosylated amino acids, sulfated amino acids, prenylated (e.g., farnesylated, geranylgeranylated) amino acids, PEG modified (e.g., "PEGylated") amino acids, biotinylated amino acids, carboxylated amino acids, phosphorylated amino acids, etc. References adequate to guide one of skill in the modification of amino acids are replete throughout the literature.

IDENTIFICATION OF ADDITIONAL FACTORS

A transcription factor provided by the present invention can also be used to identify additional endogenous or exogenous molecules that can affect a phenotype or trait of

interest. On the one hand, such molecules include organic (small or large molecules) and/or inorganic compounds that affect expression of (i.e., regulate) a particular transcription factor. Alternatively, such molecules include endogenous molecules that are acted upon either at a transcriptional level by a transcription factor of the invention to modify a phenotype as desired.

5 For example, the transcription factors can be employed to identify one or more downstream gene with which is subject to a regulatory effect of the transcription factor. In one approach, a transcription factor or transcription factor homologue of the invention is expressed in a host cell, e.g., a transgenic plant cell, tissue or explant, and expression products, either RNA or protein, of likely or random targets are monitored, e.g., by hybridization to a microarray of nucleic acid

10 probes corresponding to genes expressed in a tissue or cell type of interest, by two-dimensional gel electrophoresis of protein products, or by any other method known in the art for assessing expression of gene products at the level of RNA or protein. Alternatively, a transcription factor of the invention can be used to identify promoter sequences (i.e., binding sites) involved in the regulation of a downstream target. After identifying a promoter sequence, interactions between

15 the transcription factor and the promoter sequence can be modified by changing specific nucleotides in the promoter sequence or specific amino acids in the transcription factor that interact with the promoter sequence to alter a plant trait. Typically, transcription factor DNA binding sites are identified by gel shift assays. After identifying the promoter regions, the promoter region sequences can be employed in double-stranded DNA arrays to identify

20 molecules that affect the interactions of the transcription factors with their promoters (Bulyk et al. (1999) Nature Biotechnology 17:573-577).

The identified transcription factors are also useful to identify proteins that modify the activity of the transcription factor. Such modification can occur by covalent modification, such as by phosphorylation, or by protein-protein (homo or-heteropolymer) interactions. Any 25 method suitable for detecting protein-protein interactions can be employed. Among the methods that can be employed are co-immunoprecipitation, cross-linking and co-purification through gradients or chromatographic columns, and the two-hybrid yeast system.

The two-hybrid system detects protein interactions *in vivo* and is described in Chien, et al., (1991), Proc. Natl. Acad. Sci. USA 88, 9578-9582 and is commercially available 30 from Clontech (Palo Alto, Calif.). In such a system, plasmids are constructed that encode two hybrid proteins: one consists of the DNA-binding domain of a transcription activator protein fused to the TF polypeptide and the other consists of the transcription activator protein's activation domain fused to an unknown protein that is encoded by a cDNA that has been recombined into the plasmid as part of a cDNA library. The DNA-binding domain fusion plasmid

and the cDNA library are transformed into a strain of the yeast *Saccharomyces cerevisiae* that contains a reporter gene (e.g., lacZ) whose regulatory region contains the transcription activator's binding site. Either hybrid protein alone cannot activate transcription of the reporter gene. Interaction of the two hybrid proteins reconstitutes the functional activator protein and results in expression of the reporter gene, which is detected by an assay for the reporter gene product. Then, the library plasmids responsible for reporter gene expression are isolated and sequenced to identify the proteins encoded by the library plasmids. After identifying proteins that interact with the transcription factors, assays for compounds that interfere with the TF protein-protein interactions can be preformed.

10 **IDENTIFICATION OF MODULATORS**

In addition to the intracellular molecules described above, extracellular molecules that alter activity or expression of a transcription factor, either directly or indirectly, can be identified. For example, the methods can entail first placing a candidate molecule in contact with a plant or plant cell. The molecule can be introduced by topical administration, such as spraying or soaking of a plant, and then the molecule's effect on the expression or activity of the TF polypeptide or the expression of the polynucleotide monitored. Changes in the expression of the TF polypeptide can be monitored by use of polyclonal or monoclonal antibodies, gel electrophoresis or the like. Changes in the expression of the corresponding polynucleotide sequence can be detected by use of microarrays, Northerns, quantitative PCR, or any other technique for monitoring changes in mRNA expression. These techniques are exemplified in Ausubel et al. (eds) *Current Protocols in Molecular Biology*, John Wiley & Sons (1998). Such changes in the expression levels can be correlated with modified plant traits and thus identified molecules can be useful for soaking or spraying on fruit, vegetable and grain crops to modify traits in plants.

25 Essentially any available composition can be tested for modulatory activity of expression or activity of any nucleic acid or polypeptide herein. Thus, available libraries of compounds such as chemicals, polypeptides, nucleic acids and the like can be tested for modulatory activity. Often, potential modulator compounds can be dissolved in aqueous or organic (e.g., DMSO-based) solutions for easy delivery to the cell or plant of interest in which the 30 activity of the modulator is to be tested. Optionally, the assays are designed to screen large modulator composition libraries by automating the assay steps and providing compounds from any convenient source to assays, which are typically run in parallel (e.g., in microtiter formats on microtiter plates in robotic assays).

In one embodiment, high throughput screening methods involve providing a combinatorial library containing a large number of potential compounds (potential modulator compounds). Such "combinatorial chemical libraries" are then screened in one or more assays, as described herein, to identify those library members (particular chemical species or subclasses) that display a desired characteristic activity. The compounds thus identified can serve as target compounds.

A combinatorial chemical library can be, e.g., a collection of diverse chemical compounds generated by chemical synthesis or biological synthesis. For example, a combinatorial chemical library such as a polypeptide library is formed by combining a set of 10 chemical building blocks (e.g., in one example, amino acids) in every possible way for a given compound length (i.e., the number of amino acids in a polypeptide compound of a set length). Exemplary libraries include peptide libraries, nucleic acid libraries, antibody libraries (see, e.g., Vaughn et al. (1996) Nature Biotechnology, 14(3):309-314 and PCT/US96/10287), carbohydrate libraries (see, e.g., Liang et al. Science (1996) 274:1520-1522 and U.S. Patent 5,593,853), 15 peptide nucleic acid libraries (see, e.g., U.S. Patent 5,539,083), and small organic molecule libraries (see, e.g., benzodiazepines, Baum C&EN Jan 18, page 33 (1993); isoprenoids, U.S. Patent 5,569,588; thiazolidinones and metathiazanones, U.S. Patent 5,549,974; pyrrolidines, U.S. Patents 5,525,735 and 5,519,134; morpholino compounds, U.S. Patent 5,506,337) and the like.

Preparation and screening of combinatorial or other libraries is well known to 20 those of skill in the art. Such combinatorial chemical libraries include, but are not limited to, peptide libraries (see, e.g., U.S. Patent 5,010,175, Furka, Int. J. Pept. Prot. Res. 37:487-493 (1991) and Houghton et al. Nature 354:84-88 (1991)). Other chemistries for generating chemical diversity libraries can also be used.

In addition, as noted, compound screening equipment for high-throughput 25 screening is generally available, e.g., using any of a number of well known robotic systems that have also been developed for solution phase chemistries useful in assay systems. These systems include automated workstations including an automated synthesis apparatus and robotic systems utilizing robotic arms. Any of the above devices are suitable for use with the present invention, e.g., for high-throughput screening of potential modulators. The nature and implementation of 30 modifications to these devices (if any) so that they can operate as discussed herein will be apparent to persons skilled in the relevant art.

Indeed, entire high throughput screening systems are commercially available. These systems typically automate entire procedures including all sample and reagent pipetting, liquid dispensing, timed incubations, and final readings of the microplate in detector(s)

appropriate for the assay. These configurable systems provide high throughput and rapid start up as well as a high degree of flexibility and customization. Similarly, microfluidic implementations of screening are also commercially available.

The manufacturers of such systems provide detailed protocols the various high throughput. Thus, for example, Zymark Corp. provides technical bulletins describing screening systems for detecting the modulation of gene transcription, ligand binding, and the like. The integrated systems herein, in addition to providing for sequence alignment and, optionally, synthesis of relevant nucleic acids, can include such screening apparatus to identify modulators that have an effect on one or more polynucleotides or polypeptides according to the present invention.

In some assays it is desirable to have positive controls to ensure that the components of the assays are working properly. At least two types of positive controls are appropriate. That is, known transcriptional activators or inhibitors can be incubated with cells/plants/ etc. in one sample of the assay, and the resulting increase/decrease in transcription can be detected by measuring the resulting increase in RNA/ protein expression, etc., according to the methods herein. It will be appreciated that modulators can also be combined with transcriptional activators or inhibitors to find modulators which inhibit transcriptional activation or transcriptional repression. Either expression of the nucleic acids and proteins herein or any additional nucleic acids or proteins activated by the nucleic acids or proteins herein, or both, can be monitored.

In an embodiment, the invention provides a method for identifying compositions that modulate the activity or expression of a polynucleotide or polypeptide of the invention. For example, a test compound, whether a small or large molecule, is placed in contact with a cell, plant (or plant tissue or explant), or composition comprising the polynucleotide or polypeptide of interest and a resulting effect on the cell, plant, (or tissue or explant) or composition is evaluated by monitoring, either directly or indirectly, one or more of: expression level of the polynucleotide or polypeptide, activity (or modulation of the activity) of the polynucleotide or polypeptide. In some cases, an alteration in a plant phenotype can be detected following contact of a plant (or plant cell, or tissue or explant) with the putative modulator, e.g., by modulation of expression or activity of a polynucleotide or polypeptide of the invention.

SUBSEQUENCES

Also contemplated are uscs of polynucleotides, also referred to herein as oligonucleotides, typically having at least 12 bases, preferably at least 15, more preferably at least

20, 30, or 50 bases, which hybridize under at least highly stringent (or ultra-high stringent or ultra-ultra- high stringent conditions) conditions to a polynucleotide sequence described above. The polynucleotides may be used as probes, primers, sense and antisense agents, and the like, according to methods as noted *supra*.

5 Subsequences of the polynucleotides of the invention, including polynucleotide fragments and oligonucleotides are useful as nucleic acid probes and primers. An oligonucleotide suitable for use as a probe or primer is at least about 15 nucleotides in length, more often at least about 18 nucleotides, often at least about 21 nucleotides, frequently at least about 30 nucleotides, or about 40 nucleotides, or more in length. A nucleic acid probe is useful in hybridization
10 protocols, e.g., to identify additional polypeptide homologues of the invention, including protocols for microarray experiments. Primers can be annealed to a complementary target DNA strand by nucleic acid hybridization to form a hybrid between the primer and the target DNA strand, and then extended along the target DNA strand by a DNA polymerase enzyme. Primer pairs can be used for amplification of a nucleic acid sequence, c.g., by the polymerase chain
15 reaction (PCR) or other nucleic-acid amplification methods. See Sambrook and Ausubel, *supra*.

In addition, the invention includes an isolated or recombinant polypeptide including a subsequence of at least about 15 contiguous amino acids encoded by the recombinant or isolated polynucleotides of the invention. For example, such polypeptides, or domains or fragments thereof, can be used as immunogens, e.g., to produce antibodies specific for the
20 polypeptide sequence, or as probes for detecting a sequence of interest. A subsequence can range in size from about 15 amino acids in length up to and including the full length of the polypeptide.

PRODUCTION OF TRANSGENIC PLANTS

Modification of Traits

The polynucleotides of the invention are favorably employed to produce
25 transgenic plants with various traits, or characteristics, that have been modified in a desirable manner, e.g., to improve the seed characteristics of a plant. For example, alteration of expression levels or patterns (e.g., spatial or temporal expression patterns) of one or more of the transcription factors (or transcription factor homologues) of the invention, as compared with the levels of the same protein found in a wild type plant, can be used to modify a plant's traits. An illustrative
30 example of trait modification, improved seed characteristics, by altering expression levels of a particular transcription factor is described further in the Examples and the Sequence Listing.

Antisense and Cosuppression Approaches

In addition to expression of the nucleic acids of the invention as gene

replacement or plant phenotype modification nucleic acids, the nucleic acids are also useful for
sense and anti-sense suppression of expression, e.g., to down-regulate expression of a nucleic
5 acid of the invention, e.g., as a further mechanism for modulating plant phenotype. That is, the
nucleic acids of the invention, or subsequences or anti-sense sequences thereof, can be used to
block expression of naturally occurring homologous nucleic acids. A variety of sense and anti-
sense technologies are known in the art, e.g., as set forth in Lichtenstein and Nellen (1997)

Antisense Technology: A Practical Approach IRL Press at Oxford University, Oxford, England.

10 In general, sense or anti-sense sequences are introduced into a cell, where they are optionally
amplified, e.g., by transcription. Such sequences include both simple oligonucleotide sequences
and catalytic sequences such as ribozymes.

For example, a reduction or elimination of expression (i.e., a "knock-out") of a
transcription factor or transcription factor homologue polypeptide in a transgenic plant, e.g., to
15 modify a plant trait, can be obtained by introducing an antisense construct corresponding to the
polypeptide of interest as a cDNA. For antisense suppression, the transcription factor or homologue
cDNA is arranged in reverse orientation (with respect to the coding sequence) relative to the
promoter sequence in the expression vector. The introduced sequence need not be the full length
cDNA or gene, and need not be identical to the cDNA or gene found in the plant type to be
20 transformed. Typically, the antisense sequence need only be capable of hybridizing to the target
gene or RNA of interest. Thus, where the introduced sequence is of shorter length, a higher
degree of homology to the endogenous transcription factor sequence will be needed for effective
antisense suppression. While antisense sequences of various lengths can be utilized, preferably,
the introduced antisense sequence in the vector will be at least 30 nucleotides in length, and
25 improved antisense suppression will typically be observed as the length of the antisense sequence
increases. Preferably, the length of the antisense sequence in the vector will be greater than 100
nucleotides. Transcription of an antisense construct as described results in the production of
RNA molecules that are the reverse complement of mRNA molecules transcribed from the
endogenous transcription factor gene in the plant cell.

30 Suppression of endogenous transcription factor gene expression can also be
achieved using a ribozyme. Ribozymes are RNA molecules that possess highly specific
endoribonuclease activity. The production and use of ribozymes are disclosed in U.S. Patent No.
4,987,071 and U.S. Patent No. 5,543,508. Synthetic ribozyme sequences including antisense
RNAs can be used to confer RNA cleaving activity on the antisense RNA, such that endogenous

mRNA molecules that hybridize to the antisense RNA are cleaved, which in turn leads to an enhanced antisense inhibition of endogenous gene expression.

Vectors in which RNA encoded by a transcription factor or transcription factor homologue cDNA is over-expressed can also be used to obtain co-suppression of a corresponding endogenous gene, e.g., in the manner described in U.S. Patent No. 5,231,020 to Jorgensen. Such co-suppression (also termed sense suppression) does not require that the entire transcription factor cDNA be introduced into the plant cells, nor does it require that the introduced sequence be exactly identical to the endogenous transcription factor gene of interest. However, as with antisense suppression, the suppressive efficiency will be enhanced as specificity of hybridization is increased, e.g., as the introduced sequence is lengthened, and/or as the sequence similarity between the introduced sequence and the endogenous transcription factor gene is increased.

Vectors expressing an untranslatable form of the transcription factor mRNA, e.g., sequences comprising one or more stop codon, or nonsense mutation) can also be used to suppress expression of an endogenous transcription factor, thereby reducing or eliminating its activity and modifying one or more traits. Methods for producing such constructs are described in U.S. Patent No. 5,583,021. Preferably, such constructs are made by introducing a premature stop codon into the transcription factor gene. Alternatively, a plant trait can be modified by gene silencing using double-strand RNA (Sharp (1999) *Genes and Development* 13: 139-141).

Another method for abolishing the expression of a gene is by insertion mutagenesis using the T-DNA of *Agrobacterium tumefaciens*. After generating the insertion mutants, the mutants can be screened to identify those containing the insertion in a transcription factor or transcription factor homologue gene. Plants containing a single transgene insertion event at the desired gene can be crossed to generate homozygous plants for the mutation (Koncz et al. (1992) *Methods in Arabidopsis Research*, World Scientific).

Alternatively, a plant phenotype can be altered by eliminating an endogenous gene, such as a transcription factor or transcription factor homologue, e.g., by homologous recombination (Kempin et al. (1997) *Nature* 389:802).

A plant trait can also be modified by using the cre-lox system (for example, as described in US Pat. No. 5,658,772). A plant genome can be modified to include first and second lox sites that are then contacted with a Cre recombinase. If the lox sites are in the same orientation, the intervening DNA sequence between the two sites is excised. If the lox sites are in the opposite orientation, the intervening sequence is inverted.

The polynucleotides and polypeptides of this invention can also be expressed in a plant in the absence of an expression cassette by manipulating the activity or expression level of

the endogenous gene by other means. For example, by ectopically expressing a gene by T-DNA activation tagging (Ichikawa et al. (1997) *Nature* 390:698-701; Kakimoto et al. (1996) *Science* 274: 982-985). This method entails transforming a plant with a gene tag containing multiple transcriptional enhancers and once the tag has inserted into the genome, expression of a flanking gene coding sequence becomes deregulated. In another example, the transcriptional machinery in a plant can be modified so as to increase transcription levels of a polynucleotide of the invention (See, e.g., PCT Publications WO 96/06166 and WO 98/53057 which describe the modification of the DNA binding specificity of zinc finger proteins by changing particular amino acids in the DNA binding motif).

10 The transgenic plant can also include the machinery necessary for expressing or altering the activity of a polypeptide encoded by an endogenous gene, for example by altering the phosphorylation state of the polypeptide to maintain it in an activated state.

15 Transgenic plants (or plant cells, or plant explants, or plant tissues) incorporating the polynucleotides of the invention and/or expressing the polypeptides of the invention can be produced by a variety of well established techniques as described above. Following construction of a vector, most typically an expression cassette, including a polynucleotide, e.g., encoding a transcription factor or transcription factor homologue, of the invention, standard techniques can be used to introduce the polynucleotide into a plant, a plant cell, a plant explant or a plant tissue of interest. Optionally, the plant cell, explant or tissue can be regenerated to produce a transgenic 20 plant.

25 The plant can be any higher plant, including gymnosperms, monocotyledonous and dicotyledonous plants. Suitable protocols are available for *Leguminosae* (alfalfa, soybean, clover, etc.), *Umbelliferae* (carrot, celery, parsnip), *Cruciferae* (cabbage, radish, rapeseed, broccoli, etc.), *Cucurbitaceae* (melons and cucumber), *Gramineae* (wheat, corn, rice, barley, millet, etc.), *Solanaceae* (potato, tomato, tobacco, peppers, etc.), and various other crops. See protocols described in Ammirato et al. (1984) *Handbook of Plant Cell Culture -Crop Species*. Macmillan Publ. Co. Shimamoto et al. (1989) *Nature* 338:274-276; Fromm et al. (1990) *Bio/Technology* 8:833-839; and Vasil et al. (1990) *Bio/Technology* 8:429-434.

30 Transformation and regeneration of both monocotyledonous and dicotyledonous plant cells is now routine, and the selection of the most appropriate transformation technique will be determined by the practitioner. The choice of method will vary with the type of plant to be transformed; those skilled in the art will recognize the suitability of particular methods for given plant types. Suitable methods can include, but are not limited to: electroporation of plant protoplasts; liposome-mediated transformation; polyethylene glycol (PEG) mediated

transformation; transformation using viruses; micro-injection of plant cells; micro-projectile bombardment of plant cells; vacuum infiltration; and *Agrobacterium tumeficiens* mediated transformation. Transformation means introducing a nucleotide sequence in a plant in a manner to cause stable or transient expression of the sequence.

5 Successful examples of the modification of plant characteristics by transformation with cloned sequences which serve to illustrate the current knowledge in this field of technology, and which are herein incorporated by reference, include: U.S. Patent Nos. 5,571,706; 5,677,175; 5,510,471; 5,750,386; 5,597,945; 5,589,615; 5,750,871; 5,268,526; 5,780,708; 5,538,880; 5,773,269; 5,736,369 and 5,610,042.

10 Following transformation, plants are preferably selected using a dominant selectable marker incorporated into the transformation vector. Typically, such a marker will confer antibiotic or herbicide resistance on the transformed plants, and selection of transformants can be accomplished by exposing the plants to appropriate concentrations of the antibiotic or herbicide.

15 After transformed plants are selected and grown to maturity, those plants showing a modified trait are identified. The modified trait can be any of those traits described above. Additionally, to confirm that the modified trait is due to changes in expression levels or activity of the polypeptide or polynucleotide of the invention can be determined by analyzing mRNA expression using Northern blots, RT-PCR or microarrays, or protein expression using 20 immunoblots or Western blots or gel shift assays.

INTEGRATED SYSTEMS—SEQUENCE IDENTITY

Additionally, the present invention may be an integrated system, computer or computer readable medium that comprises an instruction set for determining the identity of one or more sequences in a database. In addition, the instruction set can be used to generate or identify 25 sequences that meet any specified criteria. Furthermore, the instruction set may be used to associate or link certain functional benefits, such improved seed characteristics, with one or more identified sequence.

For example, the instruction set can include, e.g., a sequence comparison or other alignment program, e.g., an available program such as, for example, the Wisconsin Package 30 Version 10.0, such as BLAST, FASTA, PILEUP, FINDPATTERNS or the like (GCG, Madison, WI). Public sequence databases such as GenBank, EMBL, Swiss-Prot and PIR or private sequence databases such as PhytoSeq (Incyte Pharmaceuticals, Palo Alto, CA) can be searched.

Alignment of sequences for comparison can be conducted by the local homology algorithm of Smith and Waterman (1981) Adv. Appl. Math. 2:482, by the homology alignment algorithm of Needleman and Wunsch (1970) J. Mol. Biol. 48:443, by the search for similarity method of Pearson and Lipman (1988) Proc. Natl. Acad. Sci. U.S.A. 85: 2444, by computerized implementations of these algorithms. After alignment, sequence comparisons between two (or more) polynucleotides or polypeptides are typically performed by comparing sequences of the two sequences over a comparison window to identify and compare local regions of sequence similarity. The comparison window can be a segment of at least about 20 contiguous positions, usually about 50 to about 200, more usually about 100 to about 150 contiguous positions. A 5 description of the method is provided in Ausubel et al., *supra*.

10

A variety of methods of determining sequence relationships can be used, including manual alignment and computer assisted sequence alignment and analysis. This latter approach is a preferred approach in the present invention, due to the increased throughput afforded by computer assisted methods. As noted above, a variety of computer programs for 15 performing sequence alignment are available, or can be produced by one of skill.

One example algorithm that is suitable for determining percent sequence identity and sequence similarity is the BLAST algorithm, which is described in Altschul et al. J. Mol. Biol. 215:403-410 (1990). Software for performing BLAST analyses is publicly available, e.g., through the National Center for Biotechnology Information (<http://www.ncbi.nlm.nih.gov/>). This 20 algorithm involves first identifying high scoring sequence pairs (HSPs) by identifying short words of length W in the query sequence, which either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighborhood word score threshold (Altschul et al., *supra*). These initial neighborhood word hits act as seeds for initiating searches to find longer HSPs containing them. 25 The word hits are then extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Cumulative scores are calculated using, for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always > 0) and N (penalty score for mismatching residues; always < 0). For amino acid sequences, a scoring matrix is used to calculate the cumulative score. Extension of the word hits in each 30 direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T, and X determine the sensitivity and speed of the alignment. The BLASTN program (for nucleotide sequences) uses as defaults a wordlength (W) of 11, an

expectation (E) of 10, a cutoff of 100, M=5, N=-4, and a comparison of both strands. For amino acid sequences, the BLASTP program uses as defaults a wordlength (W) of 3, an expectation (E) of 10, and the BLOSUM62 scoring matrix (*see Henikoff & Henikoff (1989) Proc. Natl. Acad. Sci. USA 89:10915*).

5 In addition to calculating percent sequence identity, the BLAST algorithm also performs a statistical analysis of the similarity between two sequences (*see, e.g., Karlin & Altschul (1993) Proc. Natl. Acad. Sci. USA 90:5873-5787*). One measure of similarity provided by the BLAST algorithm is the smallest sum probability (P(N)), which provides an indication of the probability by which a match between two nucleotide or amino acid sequences would occur
10 by chance. For example, a nucleic acid is considered similar to a reference sequence (and, therefore, in this context, homologous) if the smallest sum probability in a comparison of the test nucleic acid to the reference nucleic acid is less than about 0.1, or less than about 0.01, and or even less than about 0.001. An additional example of a useful sequence alignment algorithm is PILEUP. PILEUP creates a multiple sequence alignment from a group of related sequences using
15 progressive, pairwise alignments. The program can align, e.g., up to 300 sequences of a maximum length of 5,000 letters.

The integrated system, or computer typically includes a user input interface allowing a user to selectively view one or more sequence records corresponding to the one or more character strings, as well as an instruction set which aligns the one or more character strings with each other or with an additional character string to identify one or more region of sequence similarity. The system may include a link of one or more character strings with a particular phenotype or gene function. Typically, the system includes a user readable output element which displays an alignment produced by the alignment instruction set.
20

25 The methods of this invention can be implemented in a localized or distributed computing environment. In a distributed environment, the methods may be implemented on a single computer comprising multiple processors or on a multiplicity of computers. The computers can be linked, e.g. through a common bus, but more preferably the computer(s) are nodes on a network. The network can be a generalized or a dedicated local or wide-area network and, in certain preferred embodiments, the computers may be components of an intra-net or an internet.

30 Thus, the invention provides methods for identifying a sequence similar or homologous to one or more polynucleotides as noted herein, or one or more target polypeptides encoded by the polynucleotides, or otherwise noted herein and may include linking or associating a given plant phenotype or gene function with a sequence. In the methods, a sequence database is

provided (locally or across an inter or intra net) and a query is made against the sequence database using the relevant sequences herein and associated plant phenotypes or gene functions.

Any sequence herein can be entered into the database, before or after querying the database. This provides for both expansion of the database and, if done before the querying step, for insertion of control sequences into the database. The control sequences can be detected by the query to ensure the general integrity of both the database and the query. As noted, the query can be performed using a web browser based interface. For example, the database can be a centralized public database such as those noted herein, and the querying can be done from a remote terminal or computer across an internet or intranet.

10

EXAMPLES

The following examples are intended to illustrate but not limit the present invention.

EXAMPLE I. FULL LENGTH GENE IDENTIFICATION AND CLONING

Putative transcription factor sequences (genomic or ESTs) related to known transcription factors were identified in the *Arabidopsis thaliana* GenBank database using the tblastn sequence analysis program using default parameters and a P-value cutoff threshold of -4 or -5 or lower, depending on the length of the query sequence. Putative transcription factor sequence hits were then screened to identify those containing particular sequence strings. If the sequence hits contained such sequence strings, the sequences were confirmed as transcription factors.

Alternatively, *Arabidopsis thaliana* cDNA libraries derived from different tissues or treatments, or genomic libraries were screened to identify novel members of a transcription family using a low stringency hybridization approach. Probes were synthesized using gene specific primers in a standard PCR reaction (annealing temperature 60° C) and labeled with ³²P dCTP using the High Prime DNA Labeling Kit (Boehringer Mannheim). Purified radiolabelled probes were added to filters immersed in Church hybridization medium (0.5 M NaPO₄ pH 7.0, 7% SDS, 1 % w/v bovine serum albumin) and hybridized overnight at 60 °C with shaking. Filters were washed two times for 45 to 60 minutes with 1xSCC, 1% SDS at 60° C.

To identify additional sequence 5' or 3' of a partial cDNA sequence in a cDNA library, 5' and 3' rapid amplification of cDNA ends (RACE) was performed using the Marathon™ cDNA amplification kit (Clontech, Palo Alto, CA). Generally, the method entailed first isolating poly(A) mRNA, performing first and second strand cDNA synthesis to generate double stranded

cDNA, blunting cDNA ends, followed by ligation of the MarathonTM Adaptor to the cDNA to form a library of adaptor-ligated ds cDNA.

Gene-specific primers were designed to be used along with adaptor specific primers for both 5' and 3' RACE reactions. Nested primers, rather than single primers, were used 5 to increase PCR specificity. Using 5' and 3' RACE reactions, 5' and 3' RACE fragments were obtained, sequenced and cloned. The process can be repeated until 5' and 3' ends of the full-length gene were identified. Then the full-length cDNA was generated by PCR using primers specific to 5' and 3' ends of the gene by end-to-end PCR.

EXAMPLE II. CONSTRUCTION OF EXPRESSION VECTORS

10 The sequence was amplified from a genomic or cDNA library using primers specific to sequences upstream and downstream of the coding region. The expression vector was pMEN20 or pMEN65, which are both derived from pMON316 (Sanders et al, (1987) Nucleic Acids Research 15:1543-58) and contain the CaMV 35S promoter to express transgenes. To clone the sequence into the vector, both pMEN20 and the amplified DNA fragment were digested 15 separately with Sall and NotI restriction enzymes at 37° C for 2 hours. The digestion products were subject to electrophoresis in a 0.8% agarose gel and visualized by ethidium bromide staining. The DNA fragments containing the sequence and the linearized plasmid were excised and purified by using a Qiaquick gel extraction kit (Qiagen, CA). The fragments of interest were ligated at a ratio of 3:1 (vector to insert). Ligation reactions using T4 DNA ligase (New England 20 Biolabs, MA) were carried out at 16° C for 16 hours. The ligated DNAs were transformed into competent cells of the *E. coli* strain DH5alpha by using the heat shock method. The transformations were plated on LB plates containing 50 mg/l kanamycin (Sigma).

25 Individual colonies were grown overnight in five milliliters of LB broth containing 50 mg/l kanamycin at 37° C. Plasmid DNA was purified by using Qiaquick Mini Prep kits (Qiagen, CA).

EXAMPLE III. TRANSFORMATION OF AGROBACTERIUM WITH THE EXPRESSION VECTOR

After the plasmid vector containing the gene was constructed, the vector was used to transform *Agrobacterium tumefaciens* cells expressing the gene products. The stock of 30 *Agrobacterium tumefaciens* cells for transformation were made as described by Nagel et al. (1990) FEMS Microbiol Letts. 67: 325-328. *Agrobacterium* strain ABI was grown in 250 ml LB medium (Sigma) overnight at 28°C with shaking until an absorbance (A_{600}) of 0.5 – 1.0 was reached. Cells were harvested by centrifugation at 4,000 x g for 15 min at 4° C. Cells were then

resuspended in 250 µl chilled buffer (1 mM HEPES, pH adjusted to 7.0 with KOH). Cells were centrifuged again as described above and resuspended in 125 µl chilled buffer. Cells were then centrifuged and resuspended two more times in the same HEPES buffer as described above at a volume of 100 µl and 750 µl, respectively. Resuspended cells were then distributed into 40 µl aliquots, quickly frozen in liquid nitrogen, and stored at -80°C.

Agrobacterium cells were transformed with plasmids prepared as described above following the protocol described by Nagel et al. For each DNA construct to be transformed, 50 – 100 ng DNA (generally resuspended in 10 mM Tris-HCl, 1 mM EDTA, pH 8.0) was mixed with 40 µl of Agrobacterium cells. The DNA/cell mixture was then transferred to a chilled cuvette with a 2mm electrode gap and subject to a 2.5 kV charge dissipated at 25 µF and 200 µF using a Gene Pulser II apparatus (Bio-Rad). After electroporation, cells were immediately resuspended in 1.0 ml LB and allowed to recover without antibiotic selection for 2 – 4 hours at 28°C in a shaking incubator. After recovery, cells were plated onto selective medium of LB broth containing 100 µg/ml spectinomycin (Sigma) and incubated for 24-48 hours at 28°C. Single colonies were then picked and inoculated in fresh medium. The presence of the plasmid construct was verified by PCR amplification and sequence analysis.

EXAMPLE IV. TRANSFORMATION OF ARABIDOPSIS PLANTS WITH AGROBACTERIUM TUMEFACIENS WITH EXPRESSION VECTOR

After transformation of *Agrobacterium tumefaciens* with plasmid vectors containing the gene, single *Agrobacterium* colonies were identified, propagated, and used to transform *Arabidopsis* plants. Briefly, 500 ml cultures of LB medium containing 50 mg/l kanamycin were inoculated with the colonies and grown at 28°C with shaking for 2 days until an absorbance (A_{600}) of > 2.0 is reached. Cells were then harvested by centrifugation at 4,000 x g for 10 min, and resuspended in infiltration medium (1/2 X Murashige and Skoog salts (Sigma), 1 X Gamborg's B-5 vitamins (Sigma), 5.0% (w/v) sucrose (Sigma), 0.044 µM benzylaminopurine (Sigma), 200 µl/L Silwet L-77 (Lehle Seeds) until an absorbance (A_{600}) of 0.8 was reached.

Prior to transformation, *Arabidopsis thaliana* seeds (ecotype Columbia) were sown at a density of ~10 plants per 4" pot onto Pro-Mix BX potting medium (Hummert International) covered with fiberglass mesh (18 mm X 16 mm). Plants were grown under continuous illumination (50-75 µE/m²/sec) at 22-23°C with 65-70% relative humidity. After about 4 weeks, primary inflorescence stems (bolts) are cut off to encourage growth of multiple secondary bolts. After flowering of the mature secondary bolts, plants were prepared for transformation by removal of all siliques and opened flowers.

The pots were then immersed upside down in the mixture of *Agrobacterium* infiltration medium as described above for 30 sec, and placed on their sides to allow draining into a 1' x 2' flat surface covered with plastic wrap. After 24 h, the plastic wrap was removed and pots are turned upright. The immersion procedure was repeated one week later, for a total of two immersions per pot. Seeds were then collected from each transformation pot and analyzed following the protocol described below.

EXAMPLE V. IDENTIFICATION OF ARABIDOPSIS PRIMARY TRANSFORMANTS

Seeds collected from the transformation pots were sterilized essentially as follows. Seeds were dispersed into in a solution containing 0.1% (v/v) Triton X-100 (Sigma) and sterile H₂O and washed by shaking the suspension for 20 min. The wash solution was then drained and replaced with fresh wash solution to wash the seeds for 20 min with shaking. After removal of the second wash solution, a solution containing 0.1% (v/v) Triton X-100 and 70% ethanol (Equistar) was added to the seeds and the suspension was shaken for 5 min. After removal of the ethanol/detergent solution, a solution containing 0.1% (v/v) Triton X-100 and 30% (v/v) bleach (Clorox) was added to the seeds, and the suspension was shaken for 10 min. After removal of the bleach/detergent solution, seeds were then washed five times in sterile distilled H₂O. The seeds were stored in the last wash water at 4° C for 2 days in the dark before being plated onto antibiotic selection medium (1 X Murashige and Skoog salts (pH adjusted to 5.7 with 1M KOH), 1 X Gamborg's B-5 vitamins, 0.9% phytagar (Life Technologies), and 50 mg/l kanamycin). Seeds were germinated under continuous illumination (50-75 µE/m²/sec) at 22-23° C. After 7-10 days of growth under these conditions, kanamycin resistant primary transformants (T₁ generation) were visible and obtained. These seedlings were transferred first to fresh selection plates where the seedlings continued to grow for 3-5 more days, and then to soil (Pro-Mix BX potting medium).

Primary transformants were crossed and progeny seeds (T₂) collected; kanamycin resistant seedlings were selected and analyzed. The expression levels of the recombinant polynucleotides in the transformants varies from about a 5% expression level increase to a least a 100% expression level increase. Similar observations are made with respect to polypeptide level expression.

30

EXAMPLE VI. IDENTIFICATION OF ARABIDOPSIS PLANTS WITH TRANSCRIPTION FACTOR GENE KNOCKOUTS

The screening of insertion mutagenized *Arabidopsis* collections for null mutants in a known target gene was essentially as described in Krysan et al (1999) *Plant Cell* 11:2283-2290. Briefly, gene-specific primers, nested by 5-250 pb to each others, were designed from the 5' and 3' regions of a known target gene. Similarly, nested sets of primers were also created specific to each of the T-DNA or transposon ends (the "right" and "left" borders). All possible combinations of gene specific and T-DNA/transposon primers were used to detect by PCR an insertion event within or close to the target gene. The amplified DNA fragments were then sequenced which allows the precise determination of the T-DNA/transposon insertion point relative to the target gene. Insertion events within the coding or intervening sequence of the genes were deconvoluted from a pool comprising a plurality of insertion events to a single unique mutant plant for functional characterization. The method is described in more detail in Yu and Adam, US Application Serial No. 09/177,733 filed October 23, 1998.

EXAMPLE VII. IDENTIFICATION OF SEED CHARACTERISTICS PHENOTYPE IN OVEREXPRESSOR OR GENE KNOCKOUT PLANTS

Experiments were performed to identify those transformants or knockouts that exhibited an improved seed characteristics. For such studies, the transformants were observed by eye or biochemical assays were performed.

Among the biochemicals that were assayed were insoluble sugars, such as arabinose, fucose, galactose, mannose, rhamnose or xylose or the like; prenyl lipids, such as lutein, beta-carotene, xanthophyll-1, xanthophyll-2, chlorophylls A or B, or alpha-, delta- or gamma-tocopherol or the like; fatty acids, such as 16:0 (palmitic acid), 16:1 (palmitoleic acid), 18:0 (stearic acid), 18:1 (oleic acid), 18:2 (linoleic acid), 20:0, 18:3 (linolenic acid), 20:1 (eicosenoic acid), 20:2, 22:1 (erucic acid) or the like; waxes, such as by altering the levels of C29, C31, or C33 alkanes; sterols, such as brassicasterol, campesterol, stigmasterol, sitosterol or stigmastanol or the like, glucosinolates, protein or oil levels

Fatty acids were measured using two methods depending on whether the tissue was from leaves or seeds. For leaves, lipids were extracted and esterified with hot methanolic H₂SO₄ and partitioned into hexane from methanolic brine. For seed fatty acids, seeds were pulverized and extracted in methanol:heptane:toluene:2,2-dimethoxypropane:H₂SO₄ (39:34:20:5:2) for 90 minutes at 80°C. After cooling to room temperature the upper phase, containing the seed fatty

acid esters, was subjected to GC analysis. Fatty acid esters from both seed and leaf tissues were analyzed with a Supelco SP-2330 column.

Glucosinolates were purified from seeds or leaves by first heating the tissue at 95°C for 10 minutes. Preheated ethanol:water (50:50) is and after heating at 95°C for a further 10 minutes, 5 the extraction solvent is applied to a DEAE Sephadex column which had been previously equilibrated with 0.5 M pyridine acetate. Desulfoglucosinolates were eluted with 300 ul water and analyzed by reverse phase HPLC monitoring at 226 nm.

For wax alkanes, samples were extracted using an identical method as fatty acids and extracts were analyzed on a HP 5890 GC coupled with a 5973 MSD. Samples were 10 chromatographed on a J&W DB35 mass spectrometer (J&W Scientific).

To measure prenyl lipids levels, seeds or leaves were pulverized with 1 to 2% pyrogallol as an antioxidant. For seeds, extracted samples were filtered and a portion removed for tocopherol and carotenoid/chlorophyll analysis by HPLC. The remaining material was saponified for sterol determination. For leaves, an aliquot was removed and diluted with methanol and 15 chlorophyll A, chlorophyll B, and total carotenoids measured by spectrophotometry by determining absorbance at 665.2 nm, 652.5 nm, and 470 nm. An aliquot was removed for tocopherol and carotenoid/chlorophyll composition by HPLC using a Waters uBondapak C18 column (4.6 mm x 150 mm). The remaining methanolic solution was saponified with 10% KOH at 80°C for one hour. The samples were cooled and diluted with a mixture of methanol and 20 water. A solution of 2% methylene chloride in hexane was mixed in and the samples were centrifuged. The aqueous methanol phase was again re-extracted 2% methylene chloride in hexane and, after centrifugation, the two upper phases were combined and evaporated. 2% methylene chloride in hexane was added to the tubes and the samples were then extracted with one ml of water. The upper phase was removed, dried, and resuspended in 400 ul of 2% 25 methylene chloride in hexane and analyzed by gas chromatography using a 50 m DB-5ms (0.25 mm ID, 0.25 um phase, J&W Scientific).

Insoluble sugar levels were measured by the method essentially described by Reiter et al., Plant Journal 12:335-345. This method analyzes the neutral sugar composition of cell wall polymers found in *Arabidopsis* leaves. Soluble sugars were separated from sugar polymers by 30 extracting leaves with hot 70% ethanol. The remaining residue containing the insoluble polysaccharides was then acid hydrolyzed with allose added as an internal standard. Sugar monomers generated by the hydrolysis were then reduced to the corresponding alditols by treatment with NaBH4, then were acetylated to generate the volatile alditol acetates which were then analyzed by GC-FID. Identity of the peaks was determined by comparing the retention times

of known sugars converted to the corresponding alditol acetates with the retention times of peaks from wild-type plant extracts. Alditol acetates were analyzed on a Supelco SP-2330 capillary column (30 m x 250 um x 0.2 um) using a temperature program beginning at 180° C for 2 minutes followed by an increase to 220° C in 4 minutes. After holding at 220° C for 10 minutes, 5 the oven temperature is increased to 240° C in 2 minutes and held at this temperature for 10 minutes and brought back to room temperature.

To identify plants with alterations in total seed oil or protein content, 150mg of seeds from T2 progeny plants were subjected to analysis by Near Infrared Reflectance (NIR) using a Foss NirSystems Model 6500 with a spinning cup transport system.

10 Table 3 shows the phenotypes observed for particular overexpressor or knockout plants and provides the SEQ ID No., the internal reference code (GID), whether a knockout or overexpressor plant was analyzed and the observed phenotype.

Table 3

GIDs	Knockout (KO) or overexpressor (OE)	Phenotype observed
G214	OE	Up to 111% increase in seed lutein
G226	OE	Up to 17% increase in seed protein content
G229	OE	Up to 11% increase in seed oil, 13% decrease in seed protein
G241	OE	Up to 13% decrease in seed oil
G464	OE	Up to 12% decrease in seed oil, 25% increase in seed protein
G663	OE	Up to 16% decrease in seed oil, 14% increase in seed protein
G776	OE	Up to 31% alteration in some seed fatty acids, including
G778	OE	Up to 32% increase in seed 18:1 fatty acid
G865	OE	Up to 39% increase seed protein; 23% increase in seed oil
G869	OE	Up to 25% alteration in some seed fatty acids
G883	OE	Up to 47% decrease in seed lutein
G938	OE	Up to 115% increase in some seed fatty acids
G1328	OE	Up to 43% decrease in seed lutein
G584	OE	Larger seeds
G668	OE	Reduced seed color

15

For a particular overexpressor that shows a less beneficial seed characteristic, it may be more useful to select a plant with a decreased expression of the particular transcription factor. For a particular knockout that shows a less beneficial seed characteristic, it may be more useful to select a plant with an increased expression of the particular transcription factor.

5 EXAMPLE VIII. IDENTIFICATION OF HOMOLOGOUS SEQUENCES

Homologous sequences from *Arabidopsis* and plant species other than *Arabidopsis* were identified using database sequence search tools, such as the Basic Local Alignment Search Tool (BLAST) (Altschul et al. (1990) *J. Mol. Biol.* 215:403-410; and Altschul et al. (1997) *Nucl. Acid Res.* 25: 3389-3402). The tblastx sequence analysis programs were employed using the 10 BLOSUM-62 scoring matrix (Henikoff, S. and Henikoff, J. G. (1992) *Proc. Natl. Acad. Sci. USA* 89: 10915-10919).

Identified *Arabidopsis* homologous sequences are provided in Figure 2 and included in the Sequence Listing. The percent sequence identity among these sequences is as low as 47% sequence identity. Additionally, the entire NCBI GenBank database was filtered for sequences 15 from all plants except *Arabidopsis thaliana* by selecting all entries in the NCBI GenBank database associated with NCBI taxonomic ID 33090 (Viridiplantae; all plants) and excluding entries associated with taxonomic ID 3701 (*Arabidopsis thaliana*). These sequences were compared to sequences representing genes of SEQ IDs Nos. 1-54 on 9/26/2000 using the Washington University TBLASTX algorithm (version 2.0a19MP). For each gene of SEQ IDs 20 Nos. 1-54, individual comparisons were ordered by probability score (P-value), where the score reflects the probability that a particular alignment occurred by chance. For example, a score of 3.6e-40 is 3.6×10^{-40} . For up to ten species, the gene with the lowest P-value (and therefore the most likely homolog) is listed in Figure 3.

In addition to P-values, comparisons were also scored by percentage identity. Percentage 25 identity reflects the degree to which two segments of DNA or protein are identical over a particular length. The ranges of percent identity between the non-*Arabidopsis* genes shown in Figure 3 and the *Arabidopsis* genes in the sequence listing are: SEQ ID No. 1: 38%-89%; SEQ ID No. 3: 50%-69%; SEQ ID No. 5: 68%-93%; SEQ ID No. 7: 69%-84%; SEQ ID No. 9: 34%-60%; SEQ ID No. 11: 52%-81%; SEQ ID No. 13: 48%-81%; SEQ ID No. 15: 37%-80%; SEQ ID No. 17: 48%-83%; SEQ ID No. 19: 31%-68%; SEQ ID No. 21: 47%-90%; SEQ ID No. 23: 57%-88%; SEQ ID No. 25: 39%-79%; SEQ ID No. 27: 35%-84%; SEQ ID No. 29: 54%-89%; SEQ ID 30 No. 31: 42%-88%; SEQ ID No. 33: 41%-75%; SEQ ID No. 35: 34%-67%; SEQ ID No. 37: 72%-86%; SEQ ID No. 39: 39%-84%; SEQ ID No. 41: 40%-58%; SEQ ID No. 43: 44%-82%; SEQ ID

No. 45: 54%-68%; SEQ ID No. 47: 48%-64%; SEQ ID No. 49: 46%-88%; SEQ ID No. 51: 52%-92%; and SEQ ID No. 53: 48%-80%.

The polynucleotides and polypeptides in the Sequence Listing and the identified homologous sequences may be stored in a computer system and have associated or linked with 5 the sequences a function, such as that the polynucleotides and polypeptides are useful for modifying the seed characteristics of a plant.

All references, publications, patents and other documents herein are incorporated by reference in their entirety for all purposes. Although the invention has been described with 10 reference to the embodiments and examples above, it should be understood that various modifications can be made without departing from the spirit of the invention.

What is claimed is:

1. A transgenic plant with modified seed characteristics, which plant comprises a recombinant polynucleotide comprising a nucleotide sequence selected from the group consisting of:
 - 5 (a) a nucleotide sequence encoding a polypeptide comprising a sequence selected from SEQ ID Nos. 2N, where N=1-27, or a complementary nucleotide sequence thereof;
 - (b) a nucleotide sequence encoding a polypeptide comprising a conservatively substituted variant of a polypeptide of (a);
 - (c) a nucleotide sequence comprising a sequence selected from those of SEQ ID Nos. 2N-10, where N=1-27, or a complementary nucleotide sequence thereof;
 - (d) a nucleotide sequence comprising silent substitutions in a nucleotide sequence of (c);
 - (e) a nucleotide sequence which hybridizes under stringent conditions to a nucleotide sequence of one or more of: (a), (b), (c), or (d);
 - (f) a nucleotide sequence comprising at least 15 consecutive nucleotides of a sequence of 15 any of (a)-(e);
 - (g) a nucleotide sequence comprising a subsequence or fragment of any of (a)-(f), which subsequence or fragment encodes a polypeptide that modifies a plant's seed characteristics;
 - (h) a nucleotide sequence having at least 31% sequence identity to a nucleotide sequence of any of (a)-(g);
 - (i) a nucleotide sequence having at least 60% identity sequence identity to a nucleotide sequence of any of (a)-(g);
 - (j) a nucleotide sequence which encodes a polypeptide having at least 31% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-27;
 - 20 (k) a nucleotide sequence which encodes a polypeptide having at least 60% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-27; and
 - (l) a nucleotide sequence which encodes a polypeptide having at least 65% sequence identity to a conserved domain of a polypeptide of SEQ ID Nos. 2N, where N=1-27.
- 20 30 2. The transgenic plant of claim 1, further comprising a constitutive, inducible, or tissue-active promoter operably linked to said nucleotide sequence.
3. The transgenic plant of claim 1, wherein the plant is selected from the group consisting of: soybean, wheat, corn, potato, cotton, rice, oilseed rape, sunflower, alfalfa, sugarcane, turf,

banana, blackberry, blueberry, strawberry, raspberry, cantaloupe, carrot, cauliflower, coffee, cucumber, eggplant, grapes, honeydew, lettuce, mango, melon, onion, papaya, peas, peppers, pineapple, spinach, squash, sweet corn, tobacco, tomato, watermelon, rosaceous fruits, and vegetable brassicas.

5

4. An isolated or recombinant polynucleotide comprising a nucleotide sequence selected from the group consisting of:

- (a) a nucleotide sequence encoding a polypeptide comprising a sequence selected from SEQ ID Nos. 2N, where N=1-27, or a complementary nucleotide sequence thereof;
- 10 (b) a nucleotide sequence encoding a polypeptide comprising a conservatively substituted variant of a polypeptide of (a);
- (c) a nucleotide sequence comprising a sequence selected from those of SEQ ID Nos. 2N-1, where N=1-27, or a complementary nucleotide sequence thereof;
- 15 (d) a nucleotide sequence comprising silent substitutions in a nucleotide sequence of (c);
- (e) a nucleotide sequence which hybridizes under stringent conditions to a nucleotide sequence of one or more of: (a), (b), (c), or (d);
- (f) a nucleotide sequence comprising at least 15 consecutive nucleotides of a sequence of any of (a)-(e);
- 20 (g) a nucleotide sequence comprising a subsequence or fragment of any of (a)-(f), which subsequence or fragment encodes a polypeptide that modifies a plant's seed characteristics;
- (h) a nucleotide sequence having at least 31% sequence identity to a nucleotide sequence of any of (a)-(g);
- 25 (i) a nucleotide sequence having at least 60% identity sequence identity to a nucleotide sequence of any of (a)-(g);
- (j) a nucleotide sequence which encodes a polypeptide having at least 31% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-27;
- (k) a nucleotide sequence which encodes a polypeptide having at least 60% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-27; and
- 30 (l) a nucleotide sequence which encodes a conserved domain of a polypeptide having at least 65% sequence identity to a conserved domain of a polypeptide of SEQ ID Nos. 2N, where N=1-27.

5. The isolated or recombinant polynucleotide of claim 4, further comprising a constitutive, inducible, or tissue-active promoter operably linked to the nucleotide sequence.
6. A cloning or expression vector comprising the isolated or recombinant polynucleotide of claim 4.
7. A cell comprising the cloning or expression vector of claim 6.
8. A transgenic plant comprising the isolated or recombinant polynucleotide of claim 4.
- 10 9. A composition produced by one or more of:
 - (a) incubating one or more polynucleotide of claim 4 with a nuclease;
 - (b) incubating one or more polynucleotide of claim 4 with a restriction enzyme;
 - (c) incubating one or more polynucleotide of claim 4 with a polymerase;
 - 15 (d) incubating one or more polynucleotide of claim 4 with a polymerase and a primer;
 - (e) incubating one or more polynucleotide of claim 4 with a cloning vector, or
 - (f) incubating one or more polynucleotide of claim 4 with a cell.
10. A composition comprising two or more different polynucleotides of claim 4.
- 20 11. An isolated or recombinant polypeptide comprising a subsequence of at least about 15 contiguous amino acids encoded by the recombinant or isolated polynucleotide of claim 4.
12. A plant ectopically expressing an isolated polypeptide of claim 11.
- 25 13. A method for producing a plant having a modified seed characteristics, the method comprising altering the expression of the isolated or recombinant polynucleotide of claim 4 or the expression levels or activity of a polypeptide of claim 11 in a plant, thereby producing a modified plant, and selecting the modified plant for improved seed characteristics thereby providing the modified plant with a modified seed characteristics.
- 30 14. The method of claim 13, wherein the polynucleotide is a polynucleotide of claim 4.

15. A method of identifying a factor that is modulated by or interacts with a polypeptide encoded by a polynucleotide of claim 4, the method comprising:
- (a) expressing a polypeptide encoded by the polynucleotide in a plant; and
 - (b) identifying at least one factor that is modulated by or interacts with the polypeptide.
- 5
16. The method of claim 15, wherein the identifying is performed by detecting binding by the polypeptide to a promoter sequence, or detecting interactions between an additional protein and the polypeptide in a yeast two hybrid system.
- 10 17. The method of claim 15, wherein the identifying is performed by detecting expression of a factor by hybridization to a microarray, subtractive hybridization or differential display.
18. A method of identifying a molecule that modulates activity or expression of a polynucleotide or polypeptide of interest, the method comprising:
- 15 (a) placing the molecule in contact with a plant comprising the polynucleotide or polypeptide encoded by the polynucleotide of claim 4; and,
- (b) monitoring one or more of:
- (i) expression level of the polynucleotide in the plant;
 - (ii) expression level of the polypeptide in the plant;
 - (iii) modulation of an activity of the polypeptide in the plant; or
 - (iv) modulation of an activity of the polynucleotide in the plant.
- 20
19. An integrated system, computer or computer readable medium comprising one or more character strings corresponding to a polynucleotide of claim 4, or to a polypeptide encoded by the polynucleotide.
- 25
20. The integrated system, computer or computer readable medium of claim 19, further comprising a link between said one or more sequence strings to a modified plant seed characteristics phenotype.
- 30
21. A method of identifying a sequence similar or homologous to one or more polynucleotides of claim 4, or one or more polypeptides encoded by the polynucleotides, the method comprising:
- (a) providing a sequence database; and,

(b) querying the sequence database with one or more target sequences corresponding to the one or more polynucleotides or to the one or more polypeptides to identify one or more sequence members of the database that display sequence similarity or homology to one or more of the one or more target sequences.

5

22. The method of claim 21, wherein the querying comprises aligning one or more of the target sequences with one or more of the one or more sequence members in the sequence database.

10 23. The method of claim 21, wherein the querying comprises identifying one or more of the one or more sequence members of the database that meet a user-selected identity criteria with one or more of the target sequences.

15 24. The method of claim 21, further comprising linking the one or more of the polynucleotides of claim 4, or encoded polypeptides, to a modified plant seed characteristics phenotype.

25. A plant comprising altered expression levels of an isolated or recombinant polynucleotide of claim 4.

20

26. A plant comprising altered expression levels or the activity of an isolated or recombinant polypeptide of claim 11.

27. A plant lacking a nucleotide sequence encoding a polypeptide of claim 11.

25

Figure 1

SEQ ID No.	GID	cDNA or protein	conserved domain
1	G214	cDNA	
2	G214	protein	22-71
3	G226	cDNA	
4	G226	protein	28-78
5	G229	cDNA	
6	G229	protein	14-120
7	G241	cDNA	
8	G241	protein	14-114
9	G464	cDNA	
10	G464	protein	7-15,70-80,125-158,183-219
11	G663	cDNA	
12	G663	protein	9-111
13	G776	cDNA	
14	G776	protein	27-175
15	G778	cDNA	
16	G778	protein	220-267
17	G865	cDNA	
18	G865	protein	36-103
19	G869	cDNA	
20	G869	protein	109-177
21	G883	cDNA	
22	G883	protein	245-302
23	G938	cDNA	
24	G938	protein	96-104
25	G1328	cDNA	
26	G1328	protein	14-119
27	G584	cDNA	
28	G584	protein	401-494
29	G668	cDNA	
30	G668	protein	13-113

Figure 2

SEQ ID No.	GID	homolog	cDNA or protein	conserved domain
31	G680	homolog of G214	cDNA	
32	G680	homolog of G214	protein	24-70
33	G682	homolog of G226	cDNA	
34	G682	homolog of G226	protein	22-53
35	G225	homolog of G226	cDNA	
36	G225	homolog of G226	protein	39-76
37	G678	homolog of G229	cDNA	
38	G678	homolog of G229	protein	14-115
39	G233	homolog of G241	cDNA	
40	G233	homolog of G241	protein	14-114
41	G463	homolog of G464	cDNA	
42	G463	homolog of G464	protein	14-23, 77-88, 130-146, 194-227
43	G2422	homolog of G663	cDNA	
44	G2422	homolog of G663	protein	9-110
45	G2421	homolog of G663	cDNA	
46	G2421	homolog of G663	protein	9-110
47	G772	homolog of G776	cDNA	
48	G772	homolog of G776	protein	27-176
49	G866	homolog of G883	cDNA	
50	G866	homolog of G883	protein	43-300
51	G941	homolog of G938	cDNA	
52	G941	homolog of G938	protein	95-103
53	G198	homolog of G1328	cDNA	
54	G198	homolog of G1328	protein	14-117

Figure 3A

SEQ IDs	Gene Ids	Genbank NID	P-value	Species
1	G214	8170933	8.80E-35	<i>Lycopersicon esculentum</i>
1	G214	9205339	1.20E-27	<i>Glycine max</i>
1	G214	8577344	1.80E-23	<i>Zea mays</i>
1	G214	9119112	2.40E-18	<i>Medicago truncatula</i>
1	G214	7660673	4.80E-15	<i>Sorghum bicolor</i>
1	G214	8213273	4.40E-14	<i>Oryza sativa</i>
1	G214	3325786	4.70E-10	<i>Gossypium hirsutum</i>
1	G214	9435251	1.50E-09	<i>Hordeum vulgare</i>
1	G214	9411569	6.80E-09	<i>Triticum aestivum</i>
1	G214	7614730	3.00E-07	<i>Lotus japonicus</i>
3	G226	4396287	5.10E-15	<i>Glycine max</i>
3	G226	9410205	1.50E-05	<i>Triticum aestivum</i>
3	G226	3857004	0.11	<i>Populus tremula x Populus tremuloides</i>
3	G226	2428139	0.35	<i>Oryza sativa</i>
5	G229	7337390	5.20E-51	<i>Lycopersicon esculentum</i>
5	G229	7244424	3.90E-50	<i>Mentha x piperita</i>
5	G229	7776053	1.30E-49	<i>Lotus japonicus</i>
5	G229	2921335	4.60E-48	<i>Gossypium hirsutum</i>
5	G229	1491932	3.60E-47	<i>Zea mays</i>
5	G229	6455590	2.20E-44	<i>Glycine max</i>
5	G229	6020191	1.60E-41	<i>Pinus taeda</i>
5	G229	7765706	4.10E-41	<i>Medicago truncatula</i>
5	G229	7629167	3.20E-40	<i>Gossypium arboreum</i>
5	G229	6850206	4.30E-40	<i>Oryza sativa</i>
7	G241	6552360	2.60E-54	<i>Nicotiana tabacum</i>
7	G241	6782745	2.20E-53	<i>Oryza sativa</i>
7	G241	8097368	5.70E-53	<i>Hordeum vulgare</i>
7	G241	20560	1.80E-52	<i>Petunia x hybrida</i>
7	G241	7217727	2.70E-52	<i>Sorghum bicolor</i>
7	G241	5891408	4.60E-52	<i>Lycopersicon esculentum</i>
7	G241	5139803	7.40E-52	<i>Glycine max</i>
7	G241	7560175	4.10E-50	<i>Medicago truncatula</i>
7	G241	8381332	1.40E-44	<i>Gossypium arboreum</i>
7	G241	4886263	1.20E-42	<i>Antirrhinum majus</i>
9	G464	6527230	3.60E-31	<i>Lycopersicon esculentum</i>
9	G464	9305572	1.10E-22	<i>Sorghum bicolor</i>
9	G464	6604917	6.70E-22	<i>Medicago truncatula</i>
9	G464	5058123	2.30E-21	<i>Glycine max</i>
9	G464	3760881	1.20E-19	<i>Oryza sativa</i>
9	G464	5044476	1.20E-17	<i>Gossypium hirsutum</i>
9	G464	9412603	6.40E-15	<i>Triticum aestivum</i>
9	G464	7777277	3.20E-13	<i>Lotus japonicus</i>
9	G464	9410371	1.70E-11	<i>Hordeum vulgare</i>
9	G464	7624108	2.10E-10	<i>Gossypium arboreum</i>
11	G663	7673087	4.10E-43	<i>Petunia integrifolia</i>
11	G663	7673091	2.60E-41	<i>Petunia x hybrida</i>
11	G663	7339148	1.30E-39	<i>Lycopersicon esculentum</i>
11	G663	7673097	1.90E-36	<i>Petunia axillaris</i>
11	G663	5048991	9.90E-34	<i>Gossypium hirsutum</i>
11	G663	6455590	2.00E-31	<i>Glycine max</i>
11	G663	7560175	1.50E-27	<i>Medicago truncatula</i>
11	G663	7244424	3.20E-26	<i>Mentha x piperita</i>
11	G663	6020191	2.90E-25	<i>Pinus taeda</i>

Figure 3B

SEQ IDs	Gene Ids	Genbank NID	P-value	Species
11	G663	4138298	3.40E-25	Oryza sativa subsp. indica
13	G776	8578423	5.80E-57	Mesembryanthemum crystallinum
13	G776	7411573	2.40E-52	Lycopersicon esculentum
13	G776	9253340	5.80E-43	Solanum tuberosum
13	G776	8383411	6.00E-43	Euphorbia esula
13	G776	7565426	1.50E-39	Medicago truncatula
13	G776	6666629	2.50E-33	Glycine max
13	G776	6732155	3.60E-33	Triticum monococcum
13	G776	7502501	3.00E-32	Gossypium arboreum
13	G776	8708684	3.80E-32	Hordeum vulgare
13	G776	9307772	2.10E-31	Sorghum bicolor
15	G778	9258500	3.10E-36	Glycine max
15	G778	9211293	9.40E-21	Oryza sativa
15	G778	4380303	7.60E-08	Lycopersicon esculentum
15	G778	7718953	4.10E-07	Medicago truncatula
15	G778	7720768	6.80E-07	Lotus japonicus
15	G778	6536575	8.70E-07	Zea mays
15	G778	1668906	0.82	Citrus sinensis
17	G865	9417297	1.70E-32	Triticum aestivum
17	G865	7206394	4.90E-29	Medicago truncatula
17	G865	7796858	5.70E-27	Glycine max
17	G865	4387560	9.20E-25	Lycopersicon esculentum
17	G865	569065	1.50E-23	Oryza sativa
17	G865	7788764	4.10E-23	Lotus japonicus
17	G865	790362	8.40E-22	Nicotiana tabacum
17	G865	7528275	5.90E-21	Mesembryanthemum crystallinum
17	G865	3264766	8.80E-20	Prunus armeniaca
17	G865	8098026	2.00E-19	Hordeum vulgare
19	G869	2213784	1.30E-19	Lycopersicon esculentum
19	G869	3065894	7.30E-19	Nicotiana tabacum
19	G869	8570080	4.20E-18	Oryza sativa
19	G869	7560260	1.50E-17	Medicago truncatula
19	G869	7534890	5.20E-14	Sorghum bicolor
19	G869	6455322	1.10E-13	Glycine max
19	G869	9362061	2.70E-13	Triticum aestivum
19	G869	7788764	5.70E-13	Lotus japonicus
19	G869	7624302	2.50E-12	Gossypium arboreum
19	G869	3858036	2.80E-12	Populus balsamifera subsp. trichocarpa
21	G883	4760595	2.40E-84	Nicotiana tabacum
21	G883	4894962	3.50E-45	Avena sativa
21	G883	6719425	1.70E-36	Glycine max
21	G883	5273248	2.80E-35	Lycopersicon esculentum
21	G883	9302479	3.00E-34	Sorghum bicolor
21	G883	6799932	1.40E-31	Medicago truncatula
21	G883	5456433	4.30E-31	Zea mays
21	G883	8706346	1.40E-30	Hordeum vulgare
21	G883	8404566	2.70E-30	Oryza sativa
21	G883	1432055	2.00E-27	Petroselinum crispum
23	G938	4239844	3.10E-180	Nicotiana tabacum
23	G938	7739794	2.30E-145	Dianthus caryophyllus
23	G938	7567728	9.60E-98	Medicago truncatula
23	G938	8894549	2.70E-93	Cicer arietinum
23	G938	8104209	9.60E-90	Lycopersicon esculentum

Figure 3C

SEQ IDs	Gene Ids	Genbank NID	P-value	Species
23	G938	6462339	4.60E-79	<i>Gossypium hirsutum</i>
23	G938	9204568	1.20E-78	<i>Glycine max</i>
23	G938	7720839	1.10E-69	<i>Lotus japonicus</i>
23	G938	7324903	1.60E-52	<i>Lycopersicon pennellii</i>
23	G938	2427923	4.20E-47	<i>Oryza sativa</i>
25	G1328	4383290	5.10E-65	<i>Lycopersicon esculentum</i>
25	G1328	1946266	1.30E-58	<i>Oryza sativa</i>
25	G1328	9264503	1.40E-53	<i>Glycine max</i>
25	G1328	8381332	1.10E-52	<i>Gossypium arboreum</i>
25	G1328	9363004	3.30E-49	<i>Triticum aestivum</i>
25	G1328	7765706	1.90E-47	<i>Medicago truncatula</i>
25	G1328	20562	3.90E-47	<i>Petunia x hybrida</i>
25	G1328	5050757	4.10E-46	<i>Gossypium hirsutum</i>
25	G1328	5860031	7.80E-45	<i>Pinus taeda</i>
25	G1328	4886263	5.30E-44	<i>Antirrhinum majus</i>
27	G584	1142618	2.30E-153	<i>Phaseolus vulgaris</i>
27	G584	4321761	2.40E-128	<i>Zea mays</i>
27	G584	9280727	9.70E-122	<i>Oryza sativa</i>
27	G584	6175251	4.80E-78	<i>Lycopersicon esculentum</i>
27	G584	9193975	2.20E-59	<i>Medicago truncatula</i>
27	G584	9364538	1.40E-53	<i>Triticum aestivum</i>
27	G584	6847033	1.70E-49	<i>Glycine max</i>
27	G584	5049283	8.90E-46	<i>Gossypium hirsutum</i>
27	G584	7781217	1.00E-43	<i>Lotus japonicus</i>
27	G584	4519200	1.20E-27	<i>Perilla frutescens</i>
29	G668	8172976	9.70E-73	<i>Medicago truncatula</i>
29	G668	9252441	1.10E-70	<i>Solanum tuberosum</i>
29	G668	5897694	1.90E-66	<i>Lycopersicon esculentum</i>
29	G668	8380712	7.00E-65	<i>Gossypium arboreum</i>
29	G668	7685936	2.20E-58	<i>Glycine max</i>
29	G668	1945280	4.60E-48	<i>Oryza sativa</i>
29	G668	20562	1.10E-40	<i>Petunia x hybrida</i>
29	G668	7217727	8.20E-37	<i>Sorghum bicolor</i>
29	G668	6552360	1.90E-36	<i>Nicotiana tabacum</i>
29	G668	4886263	5.80E-36	<i>Antirrhinum majus</i>
31	G680	9258166	5.70E-36	<i>Glycine max</i>
31	G680	9255178	3.00E-29	<i>Zea mays</i>
31	G680	5274804	1.20E-27	<i>Lycopersicon esculentum</i>
31	G680	4974199	3.00E-22	<i>Oryza sativa</i>
31	G680	3325786	2.10E-21	<i>Gossypium hirsutum</i>
31	G680	9119112	1.30E-18	<i>Medicago truncatula</i>
31	G680	7660673	3.20E-17	<i>Sorghum bicolor</i>
31	G680	7243970	6.10E-16	<i>Mentha x piperita</i>
31	G680	3858093	2.10E-10	<i>Populus balsamifera subsp. trichocarpa</i>
31	G680	8845091	3.70E-10	<i>Triticum aestivum</i>
33	G682	309571	4.40E-08	<i>Zea mays</i>
33	G682	4396287	1.10E-05	<i>Glycine max</i>
33	G682	3857004	0.00051	<i>Populus tremula x Populus tremuloides</i>
33	G682	9410205	0.00085	<i>Triticum aestivum</i>
33	G682	8382118	0.0079	<i>Gossypium arboreum</i>
33	G682	2428139	0.017	<i>Oryza sativa</i>
33	G682	7339148	0.13	<i>Lycopersicon esculentum</i>
33	G682	9302672	0.32	<i>Sorghum bicolor</i>

Figure 3D

SEQ IDs	Gene Ids	Genbank NID	P-value	Species
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35	G225	309571	0.00029	<i>Zea mays</i>
35	G225	3857004	0.001	<i>Populus tremula x Populus tremuloides</i>
35	G225	9410205	0.019	<i>Triticum aestivum</i>
35	G225	9426190	0.025	<i>Triticum turgidum subsp. durum</i>
35	G225	8382118	0.046	<i>Gossypium arboreum</i>
35	G225	6782756	0.27	<i>Oryza sativa</i>
35	G225	7721017	0.4	<i>Lotus japonicus</i>
35	G225	6020136	0.47	<i>Pinus taeda</i>
35	G225	2921331	0.48	<i>Gossypium hirsutum</i>
37	G678	7244424	8.70E-50	<i>Mentha x piperita</i>
37	G678	7776053	2.70E-46	<i>Lotus japonicus</i>
37	G678	7337390	2.90E-46	<i>Lycopersicon esculentum</i>
37	G678	2921335	2.30E-43	<i>Gossypium hirsutum</i>
37	G678	6455590	8.30E-43	<i>Glycine max</i>
37	G678	1491932	1.60E-42	<i>Zea mays</i>
37	G678	5860031	4.80E-40	<i>Pinus taeda</i>
37	G678	7765706	3.20E-38	<i>Medicago truncatula</i>
37	G678	6850206	8.20E-38	<i>Oryza sativa</i>
37	G678	7217727	2.00E-37	<i>Sorghum bicolor</i>
39	G233	6552360	6.50E-66	<i>Nicotiana tabacum</i>
39	G233	20560	7.60E-65	<i>Petunia x hybrida</i>
39	G233	5139813	1.70E-58	<i>Glycine max</i>
39	G233	5891103	3.80E-58	<i>Lycopersicon esculentum</i>
39	G233	6782745	1.80E-52	<i>Oryza sativa</i>
39	G233	7560175	1.80E-51	<i>Medicago truncatula</i>
39	G233	7217727	8.30E-51	<i>Sorghum bicolor</i>
39	G233	8097368	5.80E-49	<i>Hordeum vulgare</i>
39	G233	8381332	4.60E-43	<i>Gossypium arboreum</i>
39	G233	5048991	3.50E-41	<i>Gossypium hirsutum</i>
41	G463	6527230	4.90E-36	<i>Lycopersicon esculentum</i>
41	G463	9305572	5.50E-36	<i>Sorghum bicolor</i>
41	G463	3760881	1.20E-31	<i>Oryza sativa</i>
41	G463	6604917	1.30E-23	<i>Medicago truncatula</i>
41	G463	5058123	2.50E-21	<i>Glycine max</i>
41	G463	5044476	1.10E-19	<i>Gossypium hirsutum</i>
41	G463	9412603	1.70E-17	<i>Triticum aestivum</i>
41	G463	9419394	6.00E-17	<i>Hordeum vulgare</i>
41	G463	7624108	6.20E-17	<i>Gossypium arboreum</i>
41	G463	8547152	3.20E-16	<i>Nicotiana tabacum</i>
43	G2422	7673087	9.60E-45	<i>Petunia integrifolia</i>
43	G2422	7339148	6.30E-43	<i>Lycopersicon esculentum</i>
43	G2422	7673083	7.20E-43	<i>Petunia x hybrida</i>
43	G2422	7673097	3.30E-40	<i>Petunia axillaris</i>
43	G2422	5048991	3.30E-36	<i>Gossypium hirsutum</i>
43	G2422	6455590	3.00E-33	<i>Glycine max</i>
43	G2422	6020191	3.20E-32	<i>Pinus taeda</i>
43	G2422	309571	3.60E-30	<i>Zea mays</i>
43	G2422	7560832	9.00E-30	<i>Medicago truncatula</i>
43	G2422	9363004	1.30E-29	<i>Triticum aestivum</i>
45	G2421	7673087	1.10E-46	<i>Petunia integrifolia</i>

Figure 3E

SEQ IDs	Gene Ids	Genbank NID	P-value	Species
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45	G2421	8380196	7.30E-31	<i>Gossypium arboreum</i>
45	G2421	7673095	1.90E-30	<i>Petunia axillaris</i>
45	G2421	7339148	2.80E-30	<i>Lycopersicon esculentum</i>
45	G2421	8747182	9.00E-30	<i>Medicago truncatula</i>
45	G2421	7217727	1.30E-27	<i>Sorghum bicolor</i>
45	G2421	6073050	5.50E-27	<i>Glycine max</i>
45	G2421	1101769	7.40E-27	<i>Picea mariana</i>
47	G772	8578423	4.80E-58	<i>Mesembryanthemum crystallinum</i>
47	G772	7570276	3.00E-52	<i>Medicago truncatula</i>
47	G772	7411573	1.30E-44	<i>Lycopersicon esculentum</i>
47	G772	6341483	6.30E-33	<i>Glycine max</i>
47	G772	1279639	2.00E-32	<i>Petunia x hybrida</i>
47	G772	7722907	3.50E-32	<i>Lotus japonicus</i>
47	G772	8405571	4.70E-32	<i>Hordeum vulgare</i>
47	G772	6730945	6.40E-32	<i>Oryza sativa</i>
47	G772	9302206	2.50E-31	<i>Sorghum bicolor</i>
47	G772	5047907	1.10E-30	<i>Gossypium hirsutum</i>
49	G866	4760595	3.50E-85	<i>Nicotiana tabacum</i>
49	G866	4894962	1.70E-38	<i>Avena sativa</i>
49	G866	6719425	6.60E-35	<i>Glycine max</i>
49	G866	5273248	1.10E-33	<i>Lycopersicon esculentum</i>
49	G866	9302479	7.40E-33	<i>Sorghum bicolor</i>
49	G866	6799932	3.60E-31	<i>Medicago truncatula</i>
49	G866	4886128	4.50E-31	<i>Zea mays</i>
49	G866	8404566	1.40E-29	<i>Oryza sativa</i>
49	G866	8706346	1.10E-28	<i>Hordeum vulgare</i>
49	G866	1432055	3.50E-26	<i>Petroselinum crispum</i>
51	G941	4239844	3.80E-198	<i>Nicotiana tabacum</i>
51	G941	7739794	1.20E-141	<i>Dianthus caryophyllus</i>
51	G941	7567728	7.10E-102	<i>Medicago truncatula</i>
51	G941	8104209	3.70E-97	<i>Lycopersicon esculentum</i>
51	G941	8894549	2.10E-95	<i>Cicer arietinum</i>
51	G941	5606033	1.60E-79	<i>Glycine max</i>
51	G941	6462339	4.60E-79	<i>Gossypium hirsutum</i>
51	G941	7720839	6.60E-70	<i>Lotus japonicus</i>
51	G941	7324903	1.00E-55	<i>Lycopersicon pennellii</i>
51	G941	2427923	6.90E-47	<i>Oryza sativa</i>
53	G198	4383290	3.50E-64	<i>Lycopersicon esculentum</i>
53	G198	1946266	1.10E-58	<i>Oryza sativa</i>
53	G198	9363004	5.40E-51	<i>Triticum aestivum</i>
53	G198	8381332	6.40E-51	<i>Gossypium arboreum</i>
53	G198	9264503	1.30E-50	<i>Glycine max</i>
53	G198	5050757	4.10E-46	<i>Gossypium hirsutum</i>
53	G198	20562	9.30E-46	<i>Petunia x hybrida</i>
53	G198	7765706	2.70E-45	<i>Medicago truncatula</i>
53	G198	5860031	5.40E-45	<i>Pinus taeda</i>
53	G198	4886263	7.30E-45	<i>Antirrhinum majus</i>

MBI-17 Sequence Listing.ST25
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Pilgrim, Marsha
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Gly Gly Phe Thr Ser His Pro Pro Ser Thr Phe Gly Pro Ser Cys Asp
 340 345 350

Val Glu Tyr Thr Lys Ala Ser Thr Leu Gln His Gly Ser Val Gln Ser
 355 360 365

Arg Glu Gln Glu His Ser Glu Ala Ser Lys Ala Arg Ser Ser Leu Asp
 370 375 380

Ser Glu Asp Val Glu Asn Lys Ser Lys Pro Val Cys His Glu Gln Pro
 385 390 395 400

Ser Ala Thr Pro Glu Ser Asp Ala Lys Gly Ser Asp Gly Ala Gly Asp
 405 410 415

Arg Lys Gln Val Asp Arg Ser Ser Cys Gly Ser Asn Thr Pro Ser Ser
 420 425 430

Ser Asp Asp Val Glu Ala Asp Ala Ser Glu Arg Gln Glu Asp Gly Thr
 435 440 445

Asn Gly Glu Val Lys Glu Thr Asn Glu Asp Thr Asn Lys Pro Gln Thr
 450 455 460

Ser Glu Ser Asn Ala Arg Arg Ser Arg Ile Ser Ser Asn Ile Thr Asp
 465 470 475 480

Pro Trp Lys Ser Val Ser Asp Glu Gly Arg Ile Ala Phe Gln Ala Leu
 485 490 495

Phe Ser Arg Glu Val Leu Pro Gln Ser Phe Thr Tyr Arg Glu Glu His
 500 505 510

Arg Glu Glu Glu Gln Gln Gln Glu Gln Arg Tyr Pro Met Ala Leu
 515 520 525

Asp Leu Asn Phe Thr Ala Gln Leu Thr Pro Val Asp Asp Gln Glu Glu
 530 535 540

Lys Arg Asn Thr Gly Phe Leu Gly Ile Gly Leu Asp Ala Ser Lys Leu
 545 550 555 560

Met Ser Arg Gly Arg Thr Gly Phe Lys Pro Tyr Lys Arg Cys Ser Met
 565 570 575

Glu Ala Lys Glu Ser Arg Ile Leu Asn Asn Asn Pro Ile Ile His Val
 580 585 590

Glu Gln Lys Asp Pro Lys Arg Met Arg Leu Glu Thr Gln Ala Ser Thr
 595 600 605

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 <211> 407
 <212> DNA
 <213> Arabidopsis thaliana

<220>
 <221> CDS

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<222> (10) .. (348)
 <223> G226

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ctt agg caa act aag ttc act cga tcc cga tat gac tct gaa gaa gtg
 Leu Arg Gln Thr Lys Phe Thr Arg Ser Arg Tyr Asp Ser Glu Glu Val
 15 20 25 30

agt agc atc gaa tgg gag ttt atc agt atg acc gaa caa gaa gaa gat
 Ser Ser Ile Glu Trp Glu Phe Ile Ser Met Thr Glu Gln Glu Glu Asp
 35 40 45

ctc atc tct cga atg tac aga ctt gtc ggt aat agg tgg gat tta ata
 Leu Ile Ser Arg Met Tyr Arg Leu Val Gly Asn Arg Trp Asp Leu Ile
 50 55 60

gca gga aga gtc gta gga aga aag gca aat gag att gag aga tac tgg
 Ala Gly Arg Val Val Gly Arg Lys Ala Asn Glu Ile Glu Arg Tyr Trp
 65 70 75

att atg aga aac tct gac tat ttt tct cac aaa cga cga cgt ctt aat
 Ile Met Arg Asn Ser Asp Tyr Phe Ser His Lys Arg Arg Arg Leu Asn
 80 85 90

aat tct ccc ttt ttt tct act tct cct ctt aat ctc caa gaa aat cta
 Asn Ser Pro Phe Phe Ser Thr Ser Pro Leu Asn Leu Gln Glu Asn Leu
 95 100 105 110

aaa ttg taa agaaatcaaataaaaagctt tcaatcataaa aagtagaaca
 Lys Leu

aatcttgaat gtcttcata 407

<210> 4
 <211> 112
 <212> PRT
 <213> Arabidopsis thaliana

<400> 4

Met Asp Asn Thr Asn Arg Leu Arg Arg Gly Pro Ser Leu Arg
 1 5 10 15

Gln Thr Lys Phe Thr Arg Ser Arg Tyr Asp Ser Glu Glu Val Ser Ser
 20 25 30

Ile Glu Trp Glu Phe Ile Ser Met Thr Glu Gln Glu Glu Asp Leu Ile
 35 40 45

Ser Arg Met Tyr Arg Leu Val Gly Asn Arg Trp Asp Leu Ile Ala Gly
 50 55 60

Arg Val Val Gly Arg Lys Ala Asn Glu Ile Glu Arg Tyr Trp Ile Met
 65 70 75 80

Arg Asn Ser Asp Tyr Phe Ser His Lys Arg Arg Arg Leu Asn Asn Ser
 85 90 95

Pro Phe Phe Ser Thr Ser Pro Leu Asn Leu Gln Glu Asn Leu Lys Leu
 100 105 110

MBI-17 Sequence Listing.ST25

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<213> *Arabidopsis thaliana*

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<223> G229

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 Cys Cys Glu Lys Val Gly Ile Lys Arg Gly Arg Trp Thr Ala Glu Glu
 10 15 20

gac cag att ctc tcc aac tac att caa tcc aat ggt gaa ggt tct tgg 151
 Asp Gln Ile Leu Ser Asn Tyr Ile Gln Ser Asn Gly Glu Gly Ser Trp
 25 30 35

aga tct ctc ccc aaa aat gcc gga tta aaa agg tgt gga aag agc tgt 199
 Arg Ser Leu Pro Lys Asn Ala Gly Leu Lys Arg Cys Gly Lys Ser Cys
 40 45 50

aga ttg aga tgg ata aac tat cta aga tca gac ctc aag cgt gga aac 247
Arg Leu Arg Trp Ile Asn Tyr Leu Arg Ser Asp Leu Lys Arg Gly Asn
55 60 65

ata act cca gaa gaa gaa gaa ctc gtt gtt aaa ttg cat tcc act ttg 295
 Ile Thr Pro Glu Glu Glu Leu Val Val Lys Leu His Ser Thr Leu
 70 75 80 85

gga aac agg tgg tca cta atc gcg ggt cat cta cca ggg aga aca gac 343
 Gly Asn Arg Trp Ser Leu Ile Ala Gly His Leu Pro Gly Arg Thr Asp
 90 95 100

aac gaa ata aaa aat tat tgg aac tct cat ctc agc cgt aaa ctc cac 391
 Asn Glu Ile Lys Asn Tyr Trp Asn Ser His Leu Ser Arg Lys Leu His
 105 110 115

aac ttc att agg aag cca tcc atc tct caa gac gtc tcc gcc gta atc 439
Asn Phe Ile Arg Lys Pro Ser Ile Ser Gln Asp Val Ser Ala Val Ile
120 125 130

atg gcg aac gct tct tca gcg cca ccg ccg ccg cag gca aaa cgc aga 487
Met Ala Asn Ala Ser Ser Ala Pro Pro Pro Pro Gln Ala Lys Arg Arg
135 140 145

cct ggg aga acg agt agg tcc gct atg aaa cca aaa atc cgc aga aca 535
 Leu Gly Arg Thr Ser Arg Ser Ala Met Lys Pro Lys Ile Arg Arg Thr
 150 155 160 165

aaa act cgt aaa acg aag aaa acg tct gca cca ccg gag cct aac gcc 583
 Lys Thr Arg Lys Thr Lys Thr Ser Ala Pro Pro Glu Pro Asn Ala
 170 175 180

gat gta gct ggg gct gat aaa gaa gca tta atg gtg gag tca agt gga 631
 Asp Val Ala Gly Ala Asp Lys Glu Ala Leu Met Val Glu Ser Ser Gly
 185 190 195

gcc gag gct gag cta gga cga cca tgt gac tac tat gga gat gat gat tgt 679
Ala Glu Ala Glu Leu Gly Arg Pro Cys Asp Tyr Tyr Gly Asp Asp Cys
200 205 210

aac aaa aat ctc atg agc att aat ggc gat aat gga gtt tta acg ttt 727
Asn Lys Asn Leu Met Ser Ile Asn Gly Asp Asn Gly Val Leu Thr Phe
215 220 225

gat gat gat atc atc gat ctt ttg ttg gac gag tca gat cct ggc cac 775
Asp Asp Asp Ile Ile Asp Leu Leu Leu Asp Glu Ser Asp Pro Gly His
230 235 240 245

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ttg tac aca aac aca acg tgc ggt ggt ggg gag ttg cat aac ata 823
 Leu Tyr Thr Asn Thr Cys Gly Gly Glu Leu His Asn Ile
 250 255 260

aga gac tct gaa gga gcc aga ggg ttc tcg gat act tgg aac caa ggg 871
 Arg Asp Ser Glu Gly Ala Arg Gly Phe Ser Asp Thr Trp Asn Gln Gly
 265 270 275

aat ctc gac tgt ctt ctt cag tct tgt cca tct gtg gag tcg ttt ctc 919
 Asn Leu Asp Cys Leu Leu Gln Ser Cys Pro Ser Val Glu Ser Phe Leu
 280 285 290

aac tac gac cac caa gtt aac gac gcg tcg acg gat gag ttt atc gat 967
 Asn Tyr Asp His Gln Val Asn Asp Ala Ser Thr Asp Glu Phe Ile Asp
 295 300 305

tgg gat tgt gtt tgg caa gaa ggt agt gat aat aat ctt tgg cat gag 1015
 Trp Asp Cys Val Trp Gln Glu Gly Ser Asp Asn Asn Leu Trp His Glu
 310 315 320 325

aaa gag aat ccc gac tca atg gtc tcg tgg ctt tta gac ggt gat gat 1063
 Lys Glu Asn Pro Asp Ser Met Val Ser Trp Leu Leu Asp Gly Asp Asp
 330 335 340

gag gcc acg atc ggg aat agt aat tgt gag aac ttt gga gaa ccg tta 1111
 Glu Ala Thr Ile Gly Asn Ser Asn Cys Glu Asn Phe Gly Glu Pro Leu
 345 350 355

gat cat gac gac gaa agc gct ttg gtc gct tgg ctt ctg tca tga 1156
 Asp His Asp Asp Glu Ser Ala Leu Val Ala Trp Leu Leu Ser
 360 365 370

tgatattgtat tgatccgtta tgtaatcttt tttgtgcatt cacagtttga atc 1209

<210> 6
 <211> 371
 <212> PRT
 <213> Arabidopsis thaliana

<400> 6

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Gly Glu Gly Ser Trp Arg Ser Leu Pro Lys Asn Ala Gly Leu Lys Arg
 35 40 45

Cys Gly Lys Ser Cys Arg Leu Arg Trp Ile Asn Tyr Leu Arg Ser Asp
 50 55 60

Leu Lys Arg Gly Asn Ile Thr Pro Glu Glu Glu Leu Val Val Lys
 65 70 75 80

Leu His Ser Thr Leu Gly Asn Arg Trp Ser Leu Ile Ala Gly His Leu
 85 90 95

Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Tyr Trp Asn Ser His Leu
 100 105 110

Ser Arg Lys Leu His Asn Phe Ile Arg Lys Pro Ser Ile Ser Gln Asp
 115 120 125

MBI-17 Sequence Listing ST25

Val Ser Ala Val Ile Met Ala Asn Ala Ser Ser Ala Pro Pro Pro			
130	135	140	
Gln Ala Lys Arg Arg Leu Gly Arg Thr Ser Arg Ser Ala Met Lys Pro			
145	150	155	160
Lys Ile Arg Arg Thr Lys Thr Arg Lys Thr Lys Thr Ser Ala Pro			
165	170	175	
Pro Glu Pro Asn Ala Asp Val Ala Gly Ala Asp Lys Glu Ala Leu Met			
180	185	190	
Val Glu Ser Ser Gly Ala Glu Ala Glu Leu Gly Arg Pro Cys Asp Tyr			
195	200	205	
Tyr Gly Asp Asp Cys Asn Lys Asn Leu Met Ser Ile Asn Gly Asp Asn			
210	215	220	
Gly Val Leu Thr Phe Asp Asp Asp Ile Ile Asp Leu Leu Leu Asp Glu			
225	230	235	240
Ser Asp Pro Gly His Leu Tyr Thr Asn Thr Thr Cys Gly Gly Gly			
245	250	255	
Glu Leu His Asn Ile Arg Asp Ser Glu Gly Ala Arg Gly Phe Ser Asp			
260	265	270	
Thr Trp Asn Gln Gly Asn Leu Asp Cys Leu Leu Gln Ser Cys Pro Ser			
275	280	285	
Val Glu Ser Phe Leu Asn Tyr Asp His Gln Val Asn Asp Ala Ser Thr			
290	295	300	
Asp Glu Phe Ile Asp Trp Asp Cys Val Trp Gln Glu Gly Ser Asp Asn			
305	310	315	320
Asn Leu Trp His Glu Lys Glu Asn Pro Asp Ser Met Val Ser Trp Leu			
325	330	335	
Leu Asp Gly Asp Asp Glu Ala Thr Ile Gly Asn Ser Asn Cys Glu Asn			
340	345	350	
Phe Gly Glu Pro Leu Asp His Asp Asp Glu Ser Ala Leu Val Ala Trp			
355	360	365	
Leu Leu Ser			
370			

<210> 7
<211> 1046
<212> DNA
<213> Arabidopsis thaliana

<220>
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<222> (46)...(867)
<223> G241

MBI-17 Sequence Listing.ST25

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cca tgc tgt gag aag atg ggg ttg aag aga gga cca tgg aca cct gaa 105
Pro Cys Cys Glu Lys Met Gly Leu Lys Arg Gly Pro Trp Thr Pro Glu
5 10 15 20
gaa gat caa atc ttg gtc tct ttt atc ctc aac cat gga cat agt aac 153
Glu Asp Gln Ile Leu Val Ser Phe Ile Leu Asn His Gly His Ser Asn
25 30 35
tgg cga gcc ctc cct aag caa gct ggt ctt ttg aga tgt gga aaa agc 201
Trp Arg Ala Leu Pro Lys Gln Ala Gly Leu Leu Arg Cys Gly Lys Ser
40 45 50
tgt aga ctt agg tgg atg aac tat tta aag cct gat att aaa cgt ggc 249
Cys Arg Leu Arg Trp Met Asn Tyr Leu Lys Pro Asp Ile Lys Arg Gly
55 60 65
aat ttc acc aaa gaa gag gaa gat gct atc atc agc tta cac caa ata 297
Asn Phe Thr Lys Glu Glu Asp Ala Ile Ile Ser Leu His Gln Ile
70 75 80
ctt ggc aat aga tgg tca gcg att gca gca aaa ctg cct gga aga acc 345
Leu Gly Asn Arg Trp Ser Ala Ile Ala Ala Lys Leu Pro Gly Arg Thr
85 90 95 100
gat aac gag atc aag aac gta tgg cac act cac ttg aag aag aga ctc 393
Asp Asn Glu Ile Lys Asn Val Trp His Thr His Leu Lys Arg Leu
105 110 115
gaa gat tat caa cca gct aaa cct aag acc agc aac aaa aag aag ggt 441
Glu Asp Tyr Gln Pro Ala Lys Pro Lys Thr Ser Asn Lys Lys Gly
120 125 130
act aaa cca aaa tct gaa tcc gta ata acg agc tcg aac agt act aga 489
Thr Lys Pro Lys Ser Glu Ser Val Ile Thr Ser Ser Asn Ser Thr Arg
135 140 145
agc gaa tcg gag cta gca gat tca tca aac cct tct gga gaa agc tta 537
Ser Glu Ser Glu Leu Ala Asp Ser Ser Asn Pro Ser Gly Glu Ser Leu
150 155 160
ttt tcg aca tcg cct tcg aca agt gag gtt tct tcg atg aca ctc ata 585
Phe Ser Thr Ser Pro Ser Thr Ser Glu Val Ser Ser Met Thr Leu Ile
165 170 175 180
agc cac gac ggc tat agc aac gag att aat atg gat aac aaa ccg gga 633
Ser His Asp Gly Tyr Ser Asn Glu Ile Asn Met Asp Asn Lys Pro Gly
185 190 195
gat atc agt act atc gat caa gaa tgt gtt tct ttc gaa act ttt ggt 681
Asp Ile Ser Thr Ile Asp Gln Glu Cys Val Ser Phe Glu Thr Phe Gly
200 205 210
gcg gat atc gat gaa agc ttc tgg aaa gag aca ctg tat agc caa gat 729
Ala Asp Ile Asp Glu Ser Phe Trp Lys Glu Thr Leu Tyr Ser Gln Asp
215 220 225
gaa cac aac tac gta tcg aat gac cta gaa gtc gct ggt tta gtt gag 777
Glu His Asn Tyr Val Ser Asn Asp Leu Glu Val Ala Gly Leu Val Glu
230 235 240
ata caa caa gag ttt caa aac ttg ggc tcc gct aat aat gag atg att 825
Ile Gln Gln Glu Phe Gln Asn Leu Gly Ser Ala Asn Asn Glu Met Ile
245 250 255 260
ttt gac agt gag atg gaa ctt ctg gtt cga tgt att ggc tag 867
Phe Asp Ser Glu Met Glu Leu Leu Val Arg Cys Ile Gly
265 270
aacccggcgaaa gaacaagatc tcttagccgg gctctagttt acatgtttga ggagtaaagt 927

MBI-17 Sequence Listing ST25

gaaatggtgc aaatttagtta aggctaaagaa atccaaaagc ttttgttac cgagaaaaaa	987
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20 25 30	
Gly His Ser Asn Trp Arg Ala Leu Pro Lys Gln Ala Gly Leu Leu Arg	
35 40 45	
Cys Gly Lys Ser Cys Arg Leu Arg Trp Met Asn Tyr Leu Lys Pro Asp	
50 55 60	
Ile Lys Arg Gly Asn Phe Thr Lys Glu Glu Glu Asp Ala Ile Ile Ser	
65 70 75 80	
Leu His Gln Ile Leu Gly Asn Arg Trp Ser Ala Ile Ala Ala Lys Leu	
85 90 95	
Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Val Trp His Thr His Leu	
100 105 110	
Lys Lys Arg Leu Glu Asp Tyr Gln Pro Ala Lys Pro Lys Thr Ser Asn	
115 120 125	
Lys Lys Lys Gly Thr Lys Pro Lys Ser Glu Ser Val Ile Thr Ser Ser	
130 135 140	
Asn Ser Thr Arg Ser Glu Ser Glu Leu Ala Asp Ser Ser Asn Pro Ser	
145 150 155 160	
Gly Glu Ser Leu Phe Ser Thr Ser Pro Ser Thr Ser Glu Val Ser Ser	
165 170 175	
Met Thr Leu Ile Ser His Asp Gly Tyr Ser Asn Glu Ile Asn Met Asp	
180 185 190	
Asn Lys Pro Gly Asp Ile Ser Thr Ile Asp Gln Glu Cys Val Ser Phe	
195 200 205	
Glu Thr Phe Gly Ala Asp Ile Asp Glu Ser Phe Trp Lys Glu Thr Leu	
210 215 220	
Tyr Ser Gln Asp Glu His Asn Tyr Val Ser Asn Asp Leu Glu Val Ala	
225 230 235 240	
Gly Leu Val Glu Ile Gln Gln Glu Phe Gln Asn Leu Gly Ser Ala Asn	
245 250 255	

MBI-17 Sequence Listing.ST25

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260 265 270

Gly

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<222> (41)..(664)
<223> G464

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 Glu Leu Glu Val Gly Lys Ser Asn Leu Pro Ala Glu Ser Glu Leu Glu
 10 15 20

ttg gga tta ggg ctc agc ctc ggt ggt ggc gcg tgg aaa gag cgt ggg 151
 Leu Gly Leu Gly Leu Ser Leu Gly Gly Ala Trp Lys Glu Arg Gly
 25 30 35

agg att ctt act gct aag gat ttt cct tcc gtt ggg tct aaa cgc tct 199
 Arg Ile Leu Thr Ala Lys Asp Phe Pro Ser Val Gly Ser Lys Arg Ser
 40 45 50

gct gaa tct tcc tct cac caa gga gct tct cct cct .cg^t tca agt caa 247
 Ala Glu Ser Ser Ser His Gln Gly Ala Ser Pro Pro Arg Ser Ser Gln
 55 60 65

gtg	gta	gga	tgg	cca	cca	att	ggg	tta	cac	agg	atg	aac	agt	ttg	gtt		295
Val	Val	Gly	Trp	Pro	Pro	Ile	Gly	Leu	His	Arg	Met	Asn	Ser	Leu	Val	.	.
70				75					80					85			

aat aac caa gct atg aag gca gca aga gcg gaa gaa gga gac ggg gag .	343
Asn Asn Gln Ala Met Lys Ala Ala Arg Ala Glu Glu Gly Asp Gly Glu	
90 95 100	

aag aaa gtt gtg aag aat ggt gag ctc aaa gat gtg tca atg aag gtg 391
 Lys Lys Val Val Lys Asn Gly Glu Leu Lys Asp Val Ser Met Lys Val
 105 110 115

aat ccg aaa gtt cag ggc tta ggg ttt gtt aag gtg aat atg gat gga	439
Asn Pro Lys Val Gln Gly Leu Gly Phe Val Lys Val Asn Met Asp Gly	
120 125 130	

gtt ggt ata ggc aga aaa gtg gat atg aga gct cat tcg tct tac gaa 487
Val Gly Ile Gly Arg Lys Val Asp Met Arg Ala His Ser Ser Tyr Glu
135 140 145

aac ttg gct cag acg ctt gag gaa atg ttc ttt gga atg aca ggt act 535
Asn Leu Ala Gln Thr Leu Glu Glu Met Phe Phe Gly Met Thr Gly Thr
150 155 160 165

act tgt cga gaa acg gtt aaa cct tta agg ctt tta gat gga tca tca	583	
Thr Cys Arg Glu Thr Val Lys Pro Leu Arg Leu Leu Asp Gly Ser Ser		
170	175	180

gac ttt gta ctc act tat gaa gat aag ggg att gga tgc ttg ttg gag 631
 Asp Phe Val Leu Thr Tyr Glu Asp Lys Gly Ile Gly Cys Leu Leu Glu
 185 190 195

atg ttc cat gga gaa tgt.tta tca act cgg tga aaaggcttcg · gatcatggga 684
Met Phe His Gly Glu Cys Leu Ser Thr Arg

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200 205

acctcagaag	ctagtggact	agctccaaga	cgtcaagagc	agaaggatag	acaaagaaaac	744
aaccctgttt	agcttccctt	ccaaagctgg	cattgtttat	gtattgtttg	aggttgcaa	804
tttactcgat	actttttgaa	gaaagtattt	tggagaatat	ggataaaaagc	atgcagaagc	864
tttagatatga	tttgaatccg	gttttcggat	atggtttgc	ttaggtcatt	caattcgtag	924
ttttccagtt	tgtttcttct	ttggctgtgt	accaattatc	tatgttctgt	gagagaaaagc	984
tcttg						989

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<211> 207
<212> PRT
<213> Arabidopsis thaliana

<400> 10

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Glu	Ser	Glu	Leu	Glu	Leu	Gly	Leu	Ser	Leu	Gly	Gly	Gly	Ala		
				20			25				30				

Trp	Lys	Glu	Arg	Gly	Arg	Ile	Leu	Thr	Ala	Lys	Asp	Phe	Pro	Ser	Val
					35		40				45				

Gly	Ser	Lys	Arg	Ser	Ala	Glu	Ser	Ser	Ser	His	Gln	Gly	Ala	Ser	Pro
					50		55				60				

Pro	Arg	Ser	Ser	Gln	Val	Val	Gly	Trp	Pro	Pro	Ile	Gly	Leu	His	Arg
				65		70				75			80		

Met	Asn	Ser	Leu	Val	Asn	Asn	Gln	Ala	Met	Lys	Ala	Ala	Arg	Ala	Glu
					85				90				95		

Glu	Gly	Asp	Gly	Glu	Lys	Val	Val	Lys	Asn	Gly	Glu	Leu	Lys	Asp	
					100		105				110				

Val	Ser	Met	Lys	Val	Asn	Pro	Lys	Val	Gln	Gly	Leu	Gly	Phe	Val	Lys
					115		120				125				

Val	Asn	Met	Asp	Gly	Val	Gly	Ile	Gly	Arg	Lys	Val	Asp	Met	Arg	Ala
					130		135				140				

His	Ser	Ser	Tyr	Glu	Asn	Leu	Ala	Gln	Thr	Leu	Glu	Glu	Met	Phe	Phe
					145		150			155			160		

Gly	Met	Thr	Gly	Thr	Thr	Cys	Arg	Glu	Thr	Val	Lys	Pro	Leu	Arg	Leu
					165		170			175					

Leu	Asp	Gly	Ser	Ser	Asp	Phe	Val	Leu	Thr	Tyr	Glu	Asp	Lys	Gly	Ile
					180		185				190				

Gly	Cys	Leu	Leu	Glu	Met	Phe	His	Gly	Glu	Cys	Leu	Ser	Thr	Arg	
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MBI-17 Sequence Listing.ST25

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<213> *Arabidopsis thaliana*
<220>
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<222> (113)..(862)
<223> G663

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Met Glu		
1		
ggt tcg tcc aaa ggg ttg agg aaa ggt gca tgg act gct gaa gaa gat	166	
Gly Ser Ser Lys Gly Leu Arg Lys Gly Ala Trp Thr Ala Glu Glu Asp		
5 10 15		
agt ctc ttg agg cta tgt att gat aag tat gga gaa ggc aaa tgg cat	214	
Ser Leu Leu Arg Leu Cys Ile Asp Lys Tyr Gly Glu Gly Lys Trp His		
20 25 30		
caa gtt cct ttg aga gct ggg cta aat cga tgc aga aag agt tgt aga	262	
Gln Val Pro Leu Arg Ala Gly Leu Asn Arg Cys Arg Lys Ser Cys Arg		
35 40 45 50		
cta aga tgg ttg aac tat ttg aag cca agt atc aag aga gga aga ctt	310	
Leu Arg Trp Leu Asn Tyr Leu Lys Pro Ser Ile Lys Arg Gly Arg Leu		
55 60 65		
agc aat gat gaa gtt gat ctt ctt cgc ctt cat aag ctt cta gga	358	
Ser Asn Asp Glu Val Asp Leu Leu Arg Leu His Lys Leu Leu Gly		
70 75 80		
aat agg tgg tcc ttg att gct ggt cga ttg cct ggt cg acc gct aat	406	
Asn Arg Trp Ser Leu Ile Ala Gly Arg Leu Pro Gly Arg Thr Ala Asn		
85 90 95		
gat gtc aaa aat tac tgg aac acc cat ctg agt aaa aaa cat gag tct	454	
Asp Val Lys Asn Tyr Trp Asn Thr His Leu Ser Lys Lys His Glu Ser		
100 105 110		
tcg tgt tgt aag tct aaa atg aaa aag aaa aac att att tcc cct cct	502	
Ser Cys Cys Lys Ser Lys Met Lys Lys Asn Ile Ile Ser Pro Pro		
115 120 125 130		
aca aca ccg gtc caa aaa atc ggt gtt ttt aag cct cga cct cga tcc	550	
Thr Thr Pro Val Gln Lys Ile Gly Val Phe Lys Pro Arg Pro Arg Ser		
135 140 145		
tcc tct gtt aac aat ggt tgc agc cat ctc aat ggt ctg cca gaa gtt	598	
Phe Ser Val Asn Asn Gly Cys Ser His Leu Asn Gly Leu Pro Glu Val		
150 155 160		
gat tta att cct tca tgc ctt gga ctc aag aaa aat aat gtt tgt gaa	646	
Asp Leu Ile Pro Ser Cys Leu Gly Leu Lys Lys Asn Asn Val Cys Glu		
165 170 175		
aat agt atc aca tgt aac aaa gat gat gag aaa gat gat ttt gtg aat	694	
Asn Ser Ile Thr Cys Asn Lys Asp Asp Glu Lys Asp Asp Phe Val Asn		
180 185 190		
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Asn Leu Met Asn Gly Asp Asn Met Trp Leu Glu Asn Leu Leu Gly Glu		
195 200 205 210		
aac caa gaa gct gat gcg att gtt cct gaa gcg acg aca gct gaa cat	790	
Asn Gln Glu Ala Asp Ala Ile Val Pro Glu Ala Thr Thr Ala Glu His		
215 220 225		
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MBI-17 Sequence Listing.ST25

230

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 35 40 45

Cys Arg Leu Arg Trp Leu Asn Tyr Leu Lys Pro Ser Ile Lys Arg Gly
 50 55 60

Arg Leu Ser Asn Asp Glu Val Asp Leu Leu Leu Arg Leu His Lys Leu
 65 70 75 80

Leu Gly Asn Arg Trp Ser Leu Ile Ala Gly Arg Leu Pro Gly Arg Thr
 85 90 95

Ala Asn Asp Val Lys Asn Tyr Trp Asn Thr His Leu Ser Lys Lys His
 100 105 110

Glu Ser Ser Cys Cys Lys Ser Lys Met Lys Lys Asn Ile Ile Ser
 115 120 125

Pro Pro Thr Thr Pro Val Gln Lys Ile Gly Val Phe Lys Pro Arg Pro
 130 135 140

Arg Ser Phe Ser Val Asn Asn Gly Cys Ser His Leu Asn Gly Leu Pro
 145 150 155 160

Glu Val Asp Leu Ile Pro Ser Cys Leu Gly Leu Lys Lys Asn Asn Val
 165 170 175

Cys Glu Asn Ser Ile Thr Cys Asn Lys Asp Asp Glu Lys Asp Asp Phe
 180 185 190

Val Asn Asn Leu Met Asn Gly Asp Asn Met Trp Leu Glu Asn Leu Leu
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Gly Glu Asn Gln Glu Ala Asp Ala Ile Val Pro Glu Ala Thr Thr Ala
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MBI-17 Sequence Listing.ST25

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 Pro Gly Phe Arg Phe His Pro Thr Asp Glu Glu Leu Val Ser Tyr Tyr
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 45 50 55 60

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Glu Val Asp Ile Tyr Lys His Glu Pro Trp Asp Leu Ala Val Phe Ser
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Tyr Trp Lys Ala Thr Gly Lys Asp Arg Glu Ile Arg Arg Asp Ile Leu
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Asp Gly Leu Arg Thr Asn Trp Val Met His Glu Tyr Arg Leu Val Glu
145 150 155

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 160 165 170

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 175 180 185

tat gct ccg ttc atg gaa gag gaa tgg gct gat gat gaa gga gct ctg 687
Tyr Ala Pro Phe Met Glu.Glu Glu Trp Ala Asp Asp Glu Gly Ala Leu
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MBI-17 Sequence Listing ST25

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gaa tcg gac caa cag aat cat cat gag aat gac ctc aag ccg gag gag Glu Ser Asp Gln Gln Asn His His Glu Asn Asp Leu Lys Pro Glu Glu 255 260 265	879
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gaa act ttc aag ctt gaa atg atg agt gca gaa gct atg atc agt att Glu Thr Phe Lys Leu Glu Met Met Ser Ala Glu Ala Met Ile Ser Ile 415 420 425	1359
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MBI-17 Sequence Listing.ST25

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Phe His Pro Thr Asp Glu Glu Leu Val Ser Tyr Tyr Leu Lys Arg Lys
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Val Leu Gly Gln Pro Val Arg Phe Asp Ala Ile Gly Glu Val Asp Ile
50 55 60

Tyr Lys His Glu Pro Trp Asp Leu Ala Val Phe Ser Arg Leu Lys Thr
65 70 75 80

Arg Asp Gln Glu Trp Tyr Phe Tyr Ser Ala Leu Asp Lys Lys Tyr Gly
85 90 95

Asn Gly Ala Arg Met Asn Arg Ala Thr Asn Arg Gly Tyr Trp Lys Ala
100 105 110

Thr Gly Lys Asp Arg Glu Ile Arg Arg Asp Ile Leu Leu Leu Gly Met
115 120 125

Lys Lys Thr Leu Val Phe His Ser Gly Arg Ala Pro Asp Gly Leu Arg
130 135 140

Thr Asn Trp Val Met His Glu Tyr Arg Leu Val Glu Tyr Glu Thr Glu
145 150 155 160

Lys Asn Gly Asn Leu Val Gln Asp Ala Tyr Val Leu Cys Arg Val Phe
165 170 175

His Lys Asn Asn Ile Gly Pro Pro Ser Gly Asn Arg Tyr Ala Pro Phe
180 185 190

Met Glu Glu Glu Trp Ala Asp Asp Glu Gly Ala Leu Ile Pro Gly Ile
195 200 205

Asp Val Lys Leu Arg Leu Glu Pro Pro Val Ala Asn Gly Asn Asp
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Gln Met Asp Gln Glu Ile Gln Ser Ala Ser Lys Ser Leu Ile Asn Ile
225 230 235 240

Asn Glu Pro Pro Arg Glu Thr Ala Pro Leu Asp Ile Glu Ser Asp Gln
245 250 255

Gln Asn His His Glu Asn Asp Leu Lys Pro Glu Glu His Asn Asn Asn
260 265 270

Asn Asn Tyr Asp Glu Asn Glu Glu Thr Leu Lys Arg Glu Gln Met Glu
275 280 285

MBI-17 Sequence Listing ST25

Glu Glu Glu Arg Pro Pro Arg Pro Val Cys Val Leu Asn Lys Glu Ala
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Pro Leu Pro Leu Leu Gln Tyr Lys Arg Arg Arg Gln Ser Glu Ser Asn
 305 310 315 320

Asn Asn Ser Ser Arg Asn Thr Gln Asp His Cys Ser Ser Thr Thr Thr
 325 330 335

Thr Val Asp Asn Thr Thr Thr Leu Ile Ser Ser Ser Ala Ala Ala Thr
 340 345 350

Asn Thr Ala Ile Ser Ala Leu Leu Glu Phe Ser Leu Met Gly Ile Ser
 355 360 365

Asp Lys Lys Glu Lys Pro Gln Gln Pro Leu Arg Pro His Lys Glu Pro
 370 375 380

Leu Pro Pro Gln Thr Pro Leu Ala Ser Pro Glu Glu Lys Val Asn Asp
 385 390 395 400

Leu Gln Lys Glu Ile His Gln Met Ser Val Glu Arg Glu Thr Phe Lys
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Cys Val Pro Asn Cys His Ile Asp Asp Thr Pro Ala Ala Ala Thr Thr
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Thr Val Arg Ser Thr Thr Ala Ala Asp Ile Pro Ile Leu Asp Tyr Glu
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Val Ala Glu Leu Thr Trp Glu Asn Gly Gln Leu Gly Leu His Gly Leu
40 45 50

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55 60 65

MBI-17 Sequence Listing.ST25

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MBI-17 Sequence Listing.ST25

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His Gly Leu Gly Pro Pro Arg Val Thr Ala Ser Ser Thr Lys Tyr Ser 50 55 60			
Thr Gly Ala Gly Gly Thr Leu Glu Ser Ile Val Asp Gln Ala Thr Arg 65 70 75 80			
Leu Pro Asn Pro Lys Pro Thr Asp Glu Leu Val Pro Trp Phe His His 85 90 95			
Arg Ser Ser Arg Ala Ala Met Ala Met Asp Ala Leu Val Pro Cys Ser 100 105 110			
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Met Asp Thr Tyr Asp Val Gly Phe Thr Ser Thr Ser Met Gly Ser His 165 170 175			
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Met Glu Asp Glu Glu Lys Lys Ala Gly Gly Lys Ser Ser Val Ser 195 200 205			

MBI-17 Sequence Listing.ST25

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 245 250 255

Glu Tyr Leu Lys Gln Leu Gln Ala Gln Val Ser Met Met Ser Arg Met
 260 265 270

Asn Met Pro Ser Met Met Leu Pro Met Ala Met Gln Gln Gln Gln
 275 280 285

Leu Gln Met Ser Leu Met Ser Asn Pro Met Gly Leu Gly Met Gly Met
 290 295 300

Gly Met Pro Gly Leu Gly Leu Leu Asp Leu Asn Ser Met Asn Arg Ala
 305 310 315 320

Ala Ala Ser Ala Pro Asn Ile His Ala Asn Met Met Pro Asn Pro Phe
 325 330 335

Leu Pro Met Asn Cys Pro Ser Trp Asp Ala Ser Ser Asn Asp Ser Arg
 340 345 350

Phe Gln Ser Pro Leu Ile Pro Asp Pro Met Ser Ala Phe Leu Ala Cys
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 Met Val Ser Ala Leu
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MBI-17 Sequence Listing.ST25
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 85 90 95

Arg Val Gln Gly Pro Thr Thr Thr Thr Ile Ser His Ala Pro Arg
 100 105 110

Gly Val Ser Glu Ser Met Asn Ser Pro Pro Pro Arg Pro Gly Pro Pro
 115 120 125

Ser Thr Thr Thr Ser Trp Pro Met Thr Tyr Asn Gln Asp Ile Leu
 130 135 140

Gln Tyr Ala Gln Leu Leu Thr Ser Asn Asn Glu Val Asp Leu Ser Tyr
 145 150 155 160

Tyr Thr Ser Thr Leu Phe Ser Gln Pro Phe Ser Thr Pro Ser Ser Ser
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MBI-17 Sequence Listing, ST25

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MBI-17 Sequence Listing ST25

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Lys Ser Phe Ala Ala Ser	
320	

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<213> Arabidopsis thaliana

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Thr	Gln	Pro	Leu	Arg	Lys	Val	Arg	Ile	Ile	Val	Asn	Asp	Pro	Tyr	Ala
															45

Thr	Asp	Asp	Ser	Ser	Asp	Glu	Glu	Glu	Leu	Lys	Val	Pro	Lys	Pro	
															60

Arg	Lys	Met	Lys	Arg	Ile	Val	Arg	Glu	Ile	Asn	Phe	Pro	Ser	Met	Glu
															80

Val	Ser	Glu	Gln	Pro	Ser	Glu	Ser	Ser	Gln	Asp	Ser	Thr	Lys	Thr	
															95

Asp	Gly	Lys	Ile	Ala	Val	Ser	Ala	Ser	Pro	Ala	Val	Pro	Arg	Lys	Lys
															110

Pro	Val	Gly	Val	Arg	Gln	Arg	Lys	Trp	Gly	Lys	Trp	Ala	Ala	Glu	Ile
															125

Arg	Asp	Pro	Ile	Lys	Lys	Thr	Arg	Thr	Trp	Leu	Gly	Thr	Phe	Asp	Thr
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Leu	Glu	Glu	Ala	Ala	Lys	Ala	Tyr	Asp	Ala	Lys	Lys	Leu	Glu	Phe	Asp
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Ala	Ile	Val	Ala	Gly	Asn	Val	Ser	Thr	Lys	Arg	Asp	Val	Ser	Ser	
															175

Ser	Glu	Thr	Ser	Gln	Cys	Ser	Arg	Ser	Ser	Pro	Val	Val	Pro	Val	Glu
															190

Gln	Asp	Asp	Thr	Ser	Ala	Ser	Ala	Leu	Thr	Cys	Val	Asn	Asn	Pro	Asp
															205

Asp	Val	Ser	Thr	Val	Ala	Pro	Thr	Ala	Pro	Thr	Pro	Asn	Val	Pro	Ala
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MBI-17 Sequence Listing.ST25

Gly Gly Asn Lys Glu Thr Leu Phe Asp Phe Asp Phe Thr Asn Leu Gln
 225 230 235 240

Ile Pro Asp Phe Gly Phe Leu Ala Glu Glu Gln Gln Asp Leu Asp Phe
 245 250 255

Asp Cys Phe Leu Ala Asp Asp Gln Phe Asp Asp Phe Gly Leu Leu Asp
 260 265 270

Asp Ile Gln Gly Phe Glu Asp Asn Gly Pro Ser Ala Leu Pro Asp Phe
 275 280 285

Asp Phe Ala Asp Val Glu Asp Leu Gln Leu Ala Asp Ser Ser Phe Gly
 290 295 300

Phe Leu Asp Gln Leu Ala Pro Ile Asn Ile Ser Cys Pro Leu Lys Ser
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Phe Ala Ala Ser

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Met Ala Val Asp Leu Met Arg Phe Pro Lys Ile Asp Asp Gln
1 5 10

acg gct att cag gaa gct gca tcg caa ggt tta caa agt atg gaa cat 156
Thr Ala Ile Gln Glu Ala Ala Ser Gln Gly Leu Gln Ser Met Glu His
15 20 25 30

ctg atc cgt gtc ctc tct aac cgt ccc gaa caa caa cac aac gtt gac 204
Leu Ile Arg Val Leu Ser Asn Arg Pro Glu Gln Gln His Asn Val Asp
35 40 45

tgc tcc gag atc act gac ttc acc gtt tct aaa ttc aaa acc gtc att 252
Cys Ser Glu Ile Thr Asp Phe Thr Val Ser Lys Phe Lys Thr Val Ile
50 55 60

tct ctc ctt aac cgt act ggt cac gct cgg ttc aga cgc gga ccg gtt 300
Ser Leu Leu Asn Arg Thr Gly His Ala Arg Phe Arg Arg Gly Pro Val
65 70 75

cac tcc act tcc tct gcc gca tct cag aaa cta cag agt cag atc gtt 348
His Ser Thr Ser Ser Ala Ala Ser Gln Lys Leu Gln Ser Gln Ile Val
80 85 90

aaa aat act caa cct gag gct ccg ata gtg aga aca act acg aat cac 396
Lys Asn Thr Gln Pro Glu Ala Pro Ile Val Arg Thr Thr Asn His
95 100 110

cct caa atc gtt cct cca ccg tct agt gta aca ctc gat ttc tct aaa 444
Pro Gln Ile Val Pro Pro Pro Ser Ser Val Thr Leu Asp Phe Ser Lys
115 120 125

MBI-17 Sequence Listing ST25

cca agc atc ttc ggc acc aaa gct aag agc gcc gag ctg gaa ttc tcc Pro Ser Ile Phe Gly Thr Lys Ala Lys Ser Ala Glu Leu Glu Phe Ser 130 135 140	492
aaa gaa aac ttc agt gtt tct tta aac tcc tca ttc atg tcg tcg gcg Lys Glu Asn Phe Ser Val Ser Leu Asn Ser Ser Phe Met Ser Ser Ala 145 150 155	540
ata acc gga gac ggc agc gtc tcc aat gga aaa atc ttc ctt gct tct Ile Thr Gly Asp Gly Ser Val Ser Asn Gly Lys Ile Phe Leu Ala Ser 160 165 170	588
gct ccg tcg cag cct gtt aac tct tcc gga aaa cca ccg ttg gct ggt Ala Pro Ser Gln Pro Val Asn Ser Ser Gly Lys Pro Pro Leu Ala Gly 175 180 185 190	636
cat cct tac aga aag aga tgg ctc gag cat gag cac tca gag agt ttc His Pro Tyr Arg Lys Arg Cys Leu Glu His Glu His Ser Glu Ser Phe 195 200 205	684
tcc gga aaa gtc tcc ggc tcc qcc tac gga aag tgc cat tgc aag aaa Ser Gly Lys Val Ser Gly Ser Ala Tyr Gly Lys Cys His Cys Lys Lys 210 215 220	732
agg aaa aat cgg atg aag aga acc gtg aga gta ccg gcg ata agt gca Arg Lys Asn Arg Met Lys Arg Thr Val Arg Val Pro Ala Ile Ser Ala 225 230 235	780
aag atc gcc gat att cca ccg qac gaa tat tcg tgg agg aag tac gga Lys Ile Ala Asp Ile Pro Pro Asp Glu Tyr Ser Trp Arg Lys Tyr Gly 240 245 250	828
caa aaa ccg atc aag ggc tca cca cac cca cgt ggt tac tac aag tgc Gln Lys Pro Ile Lys Gly Ser Pro His Pro Arg Gly Tyr Tyr Lys Cys 255 260 265 270	876
agt aca ttc aga gga tgg cca gcg agg aaa cac gtt gaa cga gca tta Ser Thr Phe Arg Gly Cys Pro Ala Arg Lys His Val Glu Arg Ala Leu 275 280 285	924
gat gat cca gcg atg ctt att gtg aca tac gaa gga gag cac cgt cat Asp Asp Pro Ala Met Leu Ile Val Thr Tyr Glu Gly Glu His Arg His 290 295 300	972
aac caa tcc gcg atg cag gag aat att tct tct tca ggc att aat gat Asn Gln Ser Ala Met Gln Glu Asn Ile Ser Ser Gly Ile Asn Asp 305 310 315	1020
tta gtg ttt gcc tcg gct tga ctttttttg tactatttgt ttttgattt Leu Val Phe Ala Ser Ala 320	1071
tttgagttact ttagatggat tgaaatttgt aaattttttt attaagaaaat caatttaaat agagaaaaat tagtggtggt gaaaaaaaaaaaaaaaaaaaaaaaaaaaaaaaaaaaa aaaaa	1131 1191 1195
<210> 22 <211> 324 <212> PRT <213> Arabidopsis thaliana	
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Ile Gln Glu Ala Ala Ser Gln Gly Leu Gln Ser Met Glu His Leu Ile 20 25 30	

MBI-17 Sequence Listing ST25

Arg Val Leu Ser Asn Arg Pro Glu Gln Gln His Asn Val Asp Cys Ser
35 40 45

Glu Ile Thr Asp Phe Thr Val Ser Lys Phe Lys Thr Val Ile Ser Leu
50 55 60

Leu Asn Arg Thr Gly His Ala Arg Phe Arg Arg Gly Pro Val His Ser
65 70 75 80

Thr Ser Ser Ala Ala Ser Gln Lys Leu Gln Ser Gln Ile Val Lys Asn
85 90 95

Thr Gln Pro Glu Ala Pro Ile Val Arg Thr Thr Thr Asn His Pro Gln
100 105 110

Ile Val Pro Pro Pro Ser Ser Val Thr Leu Asp Phe Ser Lys Pro Ser
115 120 125

Ile Phe Gly Thr Lys Ala Lys Ser Ala Glu Leu Glu Phe Ser Lys Glu
130 135 140

Asn Phe Ser Val Ser Leu Asn Ser Ser Phe Met Ser Ser Ala Ile Thr
145 150 155 160

Gly Asp Gly Ser Val Ser Asn Gly Lys Ile Phe Leu Ala Ser Ala Pro
165 170 175

Ser Gln Pro Val Asn Ser Ser Gly Lys Pro Pro Leu Ala Gly His Pro
180 185 190

Tyr Arg Lys Arg Cys Leu Glu His Glu His Ser Glu Ser Phe Ser Gly
195 200 205

Lys Val Ser Gly Ser Ala Tyr Gly Lys Cys His Cys Lys Lys Arg Lys
210 215 220

Asn Arg Met Lys Arg Thr Val Arg Val Pro Ala Ile Ser Ala Lys Ile
225 230 235 240

Ala Asp Ile Pro Pro Asp Glu Tyr Ser Trp Arg Lys Tyr Gly Gln Lys
245 250 255

Pro Ile Lys Gly Ser Pro His Pro Arg Gly Tyr Tyr Lys Cys Ser Thr
260 265 270

Phe Arg Gly Cys Pro Ala Arg Lys His Val Glu Arg Ala Leu Asp Asp
275 280 285

Pro Ala Met Leu Ile Val Thr Tyr Glu Gly Glu His Arg His Asn Gln
290 295 300

Ser Ala Met Gln Glu Asn Ile Ser Ser Ser Gly Ile Asn Asp Leu Val
305 310 315 320

Phe Ala Ser Ala

MBI-17 Sequence Listing.ST25

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 tct tcc tcc aca tct ctc gat gtg tgt cca tta cca caa gct gaa caa 96
 Ser Ser Ser Thr Ser Leu Asp Val Cys Pro Leu Pro Gln Ala Glu Gln
 20 25 30

 gaa cct gta gtt gaa gat gtc gac tac acc gat gat gag atg gat gtg 144
 Glu Pro Val Val Glu Asp Val Asp Tyr Thr Asp Asp Glu Met Asp Val
 35 40 45

 gat gag ctt gag aag agg atg tgg aga gac aaa atg cgt ttg aaa cgt 192
 Asp Glu Leu Glu Lys Arg Met Trp Arg Asp Lys Met Arg Leu Lys Arg
 50 55 60

 ctc aag gag caa cag agt aag tgt aaa gaa ggc gtc gat ggt tcg aaa 240
 Leu Lys Glu Gln Gln Ser Lys Cys Lys Glu Gly Val Asp Gly Ser Lys
 65 70 75 80

 cag agg cag tcg caa gag caa gct agg agg aag aaa atg tct aga gcc 288
 Gln Arg Gln Ser Gln Glu Gln Ala Arg Arg Lys Lys Met Ser Arg Ala
 85 90 95

 caa gat ggg atc ttg aag tat atg ttg aag atg atg gaa gtt tgt aaa 336
 Gln Asp Gly Ile Leu Lys Tyr Met Leu Lys Met Met Glu Val Cys Lys
 100 105 110

 gct caa ggc ttt gtt tat ggt att att cct gag aag ggt aag cct gtg 384
 Ala Gln Gly Phe Val Tyr Gly Ile Ile Pro Glu Lys Gly Lys Pro Val
 115 120 125

 act ggt gct tcg gat aat ttg agg gaa tgg tgg aaa gat aag gtt agg 432
 Thr Gly Ala Ser Asp Asn Leu Arg Glu Trp Trp Lys Asp Lys Val Arg
 130 135 140

 ttt gat cgt aat ggt cca gct gct att gct aag tat cag tca gag aat 480
 Phe Asp Arg Asn Gly Pro Ala Ala Ile Ala Lys Tyr Gln Ser Glu Asn
 145 150 155 160

 aat att tct gga ggg agt aat gat tgt aac agc ttg gtt ggt cca aca 528
 Asn Ile Ser Gly Gly Ser Asn Asp Cys Asn Ser Leu Val Gly Pro Thr
 165 170 175

 ccg cat acg ctt cag gag ctt cag gac acg act ctt ggt tcg ctt tta 576
 Pro His Thr Leu Gln Glu Leu Gln Asp Thr Thr Leu Gly Ser Leu Leu
 180 185 190

 tcg gct ttg atg caa cat tgt gat cca ccg cag aga cgg ttt cct ttg 624
 Ser Ala Leu Met Gln His Cys Asp Pro Pro Gln Arg Arg Phe Pro Leu
 195 200 205

 gag aaa gga gtt tct cca cct tgg tgg cct aat ggg aat gaa gag tgg 672
 Glu Lys Gly Val Ser Pro Pro Trp Trp Pro Asn Gly Asn Glu Glu Trp
 210 215 220

 tgg cct cag ctt ggt tta cca aat gag caa ggt cct cct cct tat aag 720
 Trp Pro Gln Leu Gly Leu Pro Asn Glu Gln Gly Pro Pro Pro Tyr Lys
 225 230 235 240

 aag cct cat gat ttg aag aaa gct tgg aaa gtc ggt gtt tta act gcg 768
 Lys Pro His Asp Leu Lys Lys Ala Trp Lys Val Gly Val Leu Thr Ala

MBI-17 Sequence Listing ST25

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agg caa tca aaa tgc ttg cag gat aag atg acg gcg aaa gag agt gct Arg Gln Ser Lys Cys Leu Gln Asp Lys Met Thr Ala Lys Glu Ser Ala 275	280	285	864
act tgg ctt gcc att att aac caa gaa gag gtt gtg gct cgg gag ctt Thr Trp Leu Ala Ile Asn Gln Glu Glu Val Val Ala Arg Glu Leu 290	295	300	912
tat ccc gag tca tgc cct ctt tct tct tca tca tta gga agc Tyr Pro Glu Ser Cys Pro Pro Leu Ser Ser Ser Ser Leu Gly Ser 305	310	315	960
ggg tcg ctt ctc att aat gat tgt agc gag tat gac gtt gaa ggt ttc Gly Ser Leu Leu Ile Asn Asp Cys Ser Glu Tyr Asp Val Glu Gly Phe 325	330	335	1008
gag aag gaa caa cat ggt ttc gat gtg gaa gag cgg aaa cca gag ata Glu Lys Glu Gln His Gly Phe Asp Val Glu Glu Arg Lys Pro Glu Ile 340	345	350	1056
gtg atg atg cat cct cta gca agc ttt ggg gtt gct aaa atg caa cat Val Met Met His Pro Leu Ala Ser Phe Gly Val Ala Lys Met Gln His 355	360	365	1104
ttt ccc ata aag gag gag gtc gcc acc acg gta aac tta gag ttc acg Phe Pro Ile Lys Glu Glu Val Ala Thr Thr Val Asn Leu Glu Phe Thr 370	375	380	1152
aga aag agg aag cag aac aat gat atg aat gtt atg gta atg gac aga Arg Lys Arg Lys Gln Asn Asn Asp Met Asn Val Met Val Met Asp Arg 385	390	395	1200
tca gca ggt tac act tgt gag aat ggt cag tgt cct cac acg aaa atg Ser Ala Gly Tyr Thr Cys Glu Asn Gly Gln Cys Pro His Ser Lys Met 405	410	415	1248
aat ctt gga ttt caa gac agg agt tca agg gac aac cac cag atg gtt Asn Leu Gly Phe Gln Asp Arg Ser Ser Arg Asp Asn His Gln Met Val 420	425	430	1296
tgt cca tat aga gac aat cgt tta gcg tat gga gca tcc aag ttt cat Cys Pro Tyr Arg Asp Asn Arg Leu Ala Tyr Gly Ala Ser Lys Phe His 435	440	445	1344
atg ggt gga atg aaa cta gta gtt cct cag caa cca gtc caa ccg atc Met Gly Gly Met Lys Leu Val Val Pro Gln Gln Pro Val Gln Pro Ile 450	455	460	1392
gac cta tcg ggc gtt gga gtt ccg gaa aac ggg cag aag atg atc acc Asp Leu Ser Gly Val Gly Val Pro Glu Asn Gly Gln Lys Met Ile Thr 465	470	475	1440
gag ctt atg gcc atg tac gac aga aat gtc caa agc aac caa acg cct Glu Leu Met Ala Met Tyr Asp Arg Asn Val Gln Ser Asn Gln Thr Pro 485	490	495	1488
cct act ttg atg gaa aac caa agc atg gtc att gat gca aaa gca gct Pro Thr Leu Met Glu Asn Gln Ser Met Val Ile Asp Ala Lys Ala Ala 500	505	510	1536
cag aat cag cag ctg aat ttc aac agt ggc aat caa atg ttt atg caa Gln Asn Gln Gln Leu Asn Phe Asn Ser Gly Asn Gln Met Phe Met Gln 515	520	525	1584
caa ggg acg aac aac ggg gtt aac aat cgg ttc cag atg gtg ttt gat Gln Gly Thr Asn Asn Gly Val Asn Asn Arg Phe Gln Met Val Phe Asp 530	535	540	1632
tcg aca cca ttc gat atg gca gca ttc gat tac aga gat gat tgg caa			1680

MBI-17 Sequence Listing.ST25

Ser Thr Pro Phe Asp Met Ala Ala Phe Asp Tyr Arg Asp Asp Trp Gln
545 550 555 560

acc gga gca atg gaa gga atg ggg aag cag 1728
 Thr Gly Ala Met Glu Gly Met Gly Lys Gln Gln Gln Gln Gln Gln Gln
 565 570 575

cag caa gat gta tca ata tgg ttc tga 1755
Gln Gln Asp Val Ser Ile Trp Phe
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<211> 584
<212> PRT
<213> *Arabidopsis thaliana*

<400> 24

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Leu Lys Glu Gln Gln Ser Lys Cys Lys Glu Gly Val Asp Gly Ser Lys
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Gln Arg Gln Ser Gln Glu Gln Ala Arg Arg Lys Lys Met Ser Arg Ala
85 90 95

Gln Asp Gly Ile Leu Lys Tyr Met Leu Lys Met Met Glu Val Cys Lys
100 105 110

Ala Gln Gly Phe Val Tyr Gly Ile Ile Pro Glu Lys Gly Lys Pro Val
115 120 125

Thr Gly Ala Ser Asp Asn Leu Arg Glu Trp Trp Lys Asp Lys Val Arg
 130 135 140

Phe Asp Arg Asn Gly Pro Ala Ala Ile Ala Lys Tyr Gln Ser Glu Asn
 145 150 155 160

Asn Ile Ser Gly Gly Ser Asn Asp Cys Asn Ser Leu Val Gly Pro Thr
165 170 175

Pro His Thr Leu Gln Glu Leu Gln Asp Thr Thr Leu Gly Ser Leu Leu
180 185 190

Ser Ala Leu Met Gln His Cys Asp Pro Pro Gln Arg Arg Phe Pro Leu
195 200 205

Glu Lys Gly Val Ser Pro Pro Trp Trp Pro Asn Gly Asn Glu Glu Trp
210 215 220

MBI-17 Sequence Listing ST25

Trp Pro Gln Leu Gly Leu Pro Asn Glu Gln Gly Pro Pro Pro Tyr Lys
225 230 235 240

Lys Pro His Asp Leu Lys Lys Ala Trp Lys Val Gly Val Leu Thr Ala
245 250 255

Val Ile Lys His Met Ser Pro Asp Ile Ala Lys Ile Arg Lys Leu Val
260 265 270

Arg Gln Ser Lys Cys Leu Gln Asp Lys Met Thr Ala Lys Glu Ser Ala
275 280 285

Thr Trp Leu Ala Ile Ile Asn Gln Glu Glu Val Val Ala Arg Glu Leu
290 295 300

Tyr Pro Glu Ser Cys Pro Pro Leu Ser Ser Ser Ser Ser Leu Gly Ser
305 310 315 320

Gly Ser Leu Leu Ile Asn Asp Cys Ser Glu Tyr Asp Val Glu Gly Phe
325 330 335

Glu Lys Glu Gln His Gly Phe Asp Val Glu Glu Arg Lys Pro Glu Ile
340 345 350

Val Met Met His Pro Leu Ala Ser Phe Gly Val Ala Lys Met Gln His
355 360 365

Phe Pro Ile Lys Glu Glu Val Ala Thr Thr Val Asn Leu Glu Phe Thr
370 375 380

Arg Lys Arg Lys Gln Asn Asn Asp Met Asn Val Met Val Met Asp Arg
385 390 395 400

Ser Ala Gly Tyr Thr Cys Glu Asn Gly Gln Cys Pro His Ser Lys Met
405 410 415

Asn Leu Gly Phe Gln Asp Arg Ser Ser Arg Asp Asn His Gln Met Val
420 425 430

Cys Pro Tyr Arg Asp Asn Arg Leu Ala Tyr Gly Ala Ser Lys Phe His
435 440 445

Met Gly Gly Met Lys Leu Val Val Pro Gln Gln Pro Val Gln Pro Ile
450 455 460

Asp Leu Ser Gly Val Gly Val Pro Glu Asn Gly Gln Lys Met Ile Thr
465 470 475 480

Glu Leu Met Ala Met Tyr Asp Arg Asn Val Gln Ser Asn Gln Thr Pro
485 490 495

Pro Thr Leu Met Glu Asn Gln Ser Met Val Ile Asp Ala Lys Ala Ala
500 505 510

Gln Asn Gln Gln Leu Asn Phe Asn Ser Gly Asn Gln Met Phe Met Gln
515 520 525

MBI-17 Sequence Listing ST25

Gln Gly Thr Asn Asn Gly Val Asn Asn Arg Phe Gln Met Val Phe Asp
 530 535 540

Ser Thr Pro Phe Asp Met Ala Ala Phe Asp Tyr Arg Asp Asp Trp Gln
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Thr Gly Ala Met Glu Gly Met Gly Lys Gln Gln Gln Gln Gln Gln
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Gln Gln Asp Val Ser Ile Trp Phe
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<223> G1328

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 Met Gly Arg Ser Pro Cys Cys Glu Lys Lys Asn Gly Leu Lys
 1 5 10

aaa gga cca tgg act cct gag gag gat caa aag ctc att gat tat atc 156
 Lys Gly Pro Trp Thr Pro Glu Glu Asp Gln Lys Leu Ile Asp Tyr Ile
 15 20 25 30

aat ata cat ggt tat gga aat tgg aga act ctt ccc aag aat gct ggg 204
 Asn Ile His Gly Tyr Gly Asn Trp Arg Thr Leu Pro Lys Asn Ala Gly
 35 40 45

tta caa aga tgt ggt aag agt tgt cgt ctc cgg tgg acc aac tat ctc 252
 Leu Gln Arg Cys Gly Lys Ser Cys Arg Leu Arg Trp Thr Asn Tyr Leu
 50 55 60

cga cca gat att aag cgt gga aga ttc tct ttt gaa gaa gaa acc 300
 Arg Pro Asp Ile Lys Arg Gly Arg Phe Ser Phe Glu Glu Glu Thr
 65 70 75

att att caa ctt cac agc atc atg gga aac aag tgg tct gcg att gcg 348
 Ile Ile Gln Leu His Ser Ile Met Gly Asn Lys Trp Ser Ala Ile Ala
 80 85 90

gct cgt ttg cct gga aga aca gac aac gag atc aaa aac tat tgg aac 396
 Ala Arg Leu Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Tyr Trp Asn
 95 100 105 110

act cac atc aga aaa aga ctt cta aag atg gga atc gac ccg gtt aca 444
 Thr His Ile Arg Lys Arg Leu Leu Lys Met Gly Ile Asp Pro Val Thr
 115 120 125

cac act cca cgt ctt gat ctt ctc gat atc tcc tcc att ctc agc tca 492
 His Thr Pro Arg Leu Asp Leu Leu Asp Ile Ser Ser Ile Leu Ser Ser
 130 135 140

tct atc tac aac tct tcg cat cat cat cat cat cat caa caa cat 540
 Ser Ile Tyr Asn Ser Ser His His His His His His Gln Gln His
 145 150 155

atg aac atg tcg agg ctc atg agt gat ggt aat cat caa cca ttg 588
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 160 165 170

MBI-17 Sequence Listing ST25

gtt aac ccc gag ata ctc aaa ctc gca acc tct ctc ttt tca aac caa Val Asn Pro Glu Ile Leu Lys Leu Ala Thr Ser Leu Phe Ser Asn Gln 175 180 185 190	636
aac cac ccc aac aac aca cac gag aac aac acg gtt aac caa acc gaa Asn His Pro Asn Asn Thr His Glu Asn Asn Thr Val Asn Gln Thr Glu 195 200 205	684
gta aac caa tac caa acc ggt tac aac atg cct ggt aat gaa gaa tta Val Asn Gln Tyr Gln Thr Gly Tyr Asn Met Pro Gly Asn Glu Glu Leu 210 215 220	732
caa tct tgg ttc cct atc atg gat caa ttc acg aat ttc caa gac ctc Gln Ser Trp Phe Pro Ile Met Asp Gln Phe Thr Asn Phe Gln Asp Leu 225 230 235	780
atg cca atg aag acg acg gtc caa aat tca ttg tca tac gat gat gat Met Pro Met Lys Thr Val Gln Asn Ser Leu Ser Tyr Asp Asp Asp 240 245 250	828
tgt tcg aag tcc aat ttt gta tta gaa cct tat tac tcc gac ttt gct Cys Ser Lys Ser Asn Phe Val Leu Glu Pro Tyr Ser Asp Phe Ala 255 260 265 270	876
tca gtc ttg acc aca cct tct tca agc ccg act ccg tta aac tca agt Ser Val Leu Thr Thr Pro Ser Ser Pro Thr Pro Leu Asn Ser Ser 275 280 285	924
tcc tca act tac atc aat agt agc act tgc agc acc gag gat gaa aaa Ser Ser Thr Tyr Ile Asn Ser Ser Thr Cys Ser Thr Glu Asp Glu Lys 290 295 300	972
gag agt tat tac agt gat aat atc act aat tat tcg ttt gat gtt aat Glu Ser Tyr Tyr Ser Asp Asn Ile Thr Asn Tyr Ser Phe Asp Val Asn 305 310 315	1020
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Arg Cys Gly Lys Ser Cys Arg Leu Arg Trp Thr Asn Tyr Leu Arg Pro 50 55 60	
Asp Ile Lys Arg Gly Arg Phe Ser Phe Glu Glu Glu Thr Ile Ile 65 70 75 80	
Gln Leu His Ser Ile Met Gly Asn Lys Trp Ser Ala Ile Ala Arg 85 90 95	

MBI-17 Sequence Listing.ST25

Leu Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Tyr Trp Asn Thr His
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Ile Arg Lys Arg Leu Leu Lys Met Gly Ile Asp Pro Val Thr His Thr
 115 120 125

Pro Arg Leu Asp Leu Leu Asp Ile Ser Ser Ile Leu Ser Ser Ser Ile
 130 135 140

Tyr Asn Ser Ser His His His His His His Gln Gln His Met Asn
 145 150 155 160

Met Ser Arg Leu Met Met Ser Asp Gly Asn His Gln Pro Leu Val Asn
 165 170 175

Pro Glu Ile Leu Lys Leu Ala Thr Ser Leu Phe Ser Asn Gln Asn His
 180 185 190

Pro Asn Asn Thr His Glu Asn Asn Thr Val Asn Gln Thr Glu Val Asn
 195 200 205

Gln Tyr Gln Thr Gly Tyr Asn Met Pro Gly Asn Glu Glu Leu Gln Ser
 210 215 220

Trp Phe Pro Ile Met Asp Gln Phe Thr Asn Phe Gln Asp Leu Met Pro
 225 230 235 240

Met Lys Thr Thr Val Gln Asn Ser Leu Ser Tyr Asp Asp Asp Cys Ser
 245 250 255

Lys Ser Asn Phe Val Leu Glu Pro Tyr Tyr Ser Asp Phe Ala Ser Val
 260 265 270

Leu Thr Thr Pro Ser Ser Ser Pro Thr Pro Leu Asn Ser Ser Ser Ser
 275 280 285

Thr Tyr Ile Asn Ser Ser Thr Cys Ser Thr Glu Asp Glu Lys Glu Ser
 290 295 300

Tyr Tyr Ser Asp Asn Ile Thr Asn Tyr Ser Phe Asp Val Asn Gly Phe
 305 310 315 320

Leu Gln Phe Gln

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<223> G584

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Met Ser Pro Thr Asn

54

MBI-17 Sequence Listing.ST25

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40 45 50	
cct ctt cct caa gtc aac gaa gat aat ctc cag caa cgt ctc caa gct Pro Leu Pro Gln Val Asn Glu Asp Asn Leu Gln Gln Arg Leu Gln Ala	246
55 60 65	
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70 75 80 85	
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90 95 100	
ttg tta ggt tgg gga gat ggt tat tac aaa gga gaa gaa gag aag tct Leu Leu Gly Trp Gly Asp Gly Tyr Tyr Lys Gly Glu Glu Lys Ser	390
105 110 115	
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120 125 130	
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135 140 145	
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150 155 160 165	
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170 175 180	
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185 190 195	
gct tta gct gga tca agt tgt gag aga gct cgt caa ggt cag att tat Ala Leu Ala Gly Ser Ser Cys Glu Arg Ala Arg Gln Gly Gln Ile Tyr	678
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215 220 225	
ctt ggt tcg tcg gag att att cat caa agt tca gat ctt gtt gat aaa Leu Gly Ser Ser Glu Ile Ile His Gln Ser Ser Asp Leu Val Asp Lys	774
230 235 240 245	
gtt gac acc ttt ttc aat ttt aac aat ggt ggt ggt gaa ttt ggt tct Val Asp Thr Phe Asn Phe Asn Asn Gly Gly Glu Phe Gly Ser	822
250 255 260	
tgg gcg ttt aat ttg aat cca gat caa gga gag aat gat cca ggt ttg Trp Ala Phe Asn Leu Asn Pro Asp Gln Gly Glu Asn Asp Pro Gly Leu	870
265 270 275	
tgg att agt gaa cct aat ggt gtt gac tct ggt ctt gta gct gct ccg Trp Ile Ser Glu Pro Asn Gly Val Asp Ser Gly Leu Val Ala Ala Pro	918
280 285 290	
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Val Met Asn Asn Gly Gly Asn Asp Ser Thr Ser Asn Ser Asp Ser Gln 295 300 305		
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tcg aat cac tct gat ctt gaa gct tca gtg gct aaa gaa gct gag agt Ser Asn His Ser Asp Leu Glu Ala Ser Val Ala Lys Glu Ala Glu Ser 375 380 385		1206
aac aga gtt gtg gtt gaa ccg gag aag aaa ccg agg aaa cga ggg aga Asn Arg Val Val Glu Pro Glu Lys Lys Pro Arg Lys Arg Gly Arg 390 395 400 405		1254
aaa ccg gcg aat gga aga gaa gag cct ttg aat cat gta gag gca gag Lys Pro Ala Asn Gly Arg Glu Glu Pro Leu Asn His Val Glu Ala Glu 410 415 420		1302
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gtg gtt cct aat gtg tct aag atg gat aaa gct tct cta tta gga gat Val Val Pro Asn Val Ser Lys Met Asp Lys Ala Ser Leu Leu Gly Asp 440 445 450		1398
gct att tcg tat atc agt gag ctt aag tct aag ttg caa aag gct gaa Ala Ile Ser Tyr Ile Ser Glu Leu Lys Ser Lys Leu Gln Lys Ala Glu 455 460 465		1446
tct gat aaa gaa gag ttg cag aag cag att gat gtg atg aat aaa gaa Ser Asp Lys Glu Leu Gln Lys Gln Ile Asp Val Met Asn Lys Glu 470 475 480 485		1494
gcg gga aat gcg aaa agt tcg gta aaa gat cga aaa tgt ttg aat caa Ala Gly Asn Ala Lys Ser Ser Val Lys Asp Arg Lys Cys Leu Asn Gln 490 495 500		1542
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gct aag ttc atg gaa gca ctt aag gag ttg gat ttg gaa gtg aat cat Ala Lys Phe Met Glu Ala Leu Lys Glu Leu Asp Leu Glu Val Asn His 535 540 545		1686
gcg agt tta tcg gta gtg aat gat ctt atg atc caa caa gcg act gtg Ala Ser Leu Ser Val Val Asn Asp Leu Met Ile Gln Gln Ala Thr Val 550 555 560 565		1734
aaa atg ggg aat cag ttt ttc acg caa gat caa ctc aag gtt gct cta Lys Met Gly Asn Gln Phe Phe Thr Gln Asp Gln Leu Lys Val Ala Leu 570 575 580		1782
acg gag aaa gtt gga gaa tgt cca tga attgaagtca gcatcttttag Thr Glu Lys Val Gly Glu Cys Pro 585		1829

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gtgttaagggt	aattttgtag	taccacttg	ttgctattga	atgcttgttta	gagaggattc	2009
ttagtgttagt	atatgattag	gttggggtt	gttggttcat	gagataaata	aatgtgtttg	2069
atcaatggtt	aagtctttgg	tttgggttg	tatgtatgt	aataaggctt	ttgttagaaa	2129
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Met Glu Ala Phe Ile Gly Gly Ser Asp His Ser Ser Leu Phe Pro
35 40 45

Pro Leu Pro Pro Pro Pro Leu Pro Gln Val Asn Glu Asp Asn Leu Gln
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Gln Arg Leu Gln Ala Leu Ile Glu Gly Ala Asn Glu Asn Trp Thr Tyr
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Ala Val Phe Trp Gln Ser Ser His Gly Phe Ala Gly Glu Asp Asn Asn
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Asn Asn Asn Thr Val Leu Leu Gly Trp Gly Asp Gly Tyr Tyr Lys Gly
100 105 110

Glu Glu Glu Lys Ser Arg Lys Lys Ser Asn Pro Ala Ser Ala Ala
115 120 125

Glu Gln Glu His Arg Lys Arg Val Ile Arg Glu Leu Asn Ser Leu Ile
130 135 140

Ser Gly Gly Val Gly Gly Asp Glu Ala Gly Asp Glu Glu Val Thr
145 150 155 160

Asp Thr Glu Trp Phe Phe Leu Val Ser Met Thr Gln Ser Phe Val Lys
165 170 175

Gly Thr Gly Leu Pro Gly Gln Ala Phe Ser Asn Ser Asp Thr Ile Trp
180 185 190

Leu Ser Gly Ser Asn Ala Leu Ala Gly Ser Ser Cys Glu Arg Ala Arg
195 200 205

Gln Gly Gln Ile Tyr Gly Leu Gln Thr Met Val Cys Val Ala Thr Glu
210 215 220

MBI-17 Sequence Listing.ST25

Asn Gly Val Val Glu Leu Gly Ser Ser Glu Ile Ile His Gln Ser Ser
 225 230 235 240

Asp Leu Val Asp Lys Val Asp Thr Phe Phe Asn Phe Asn Asn Gly Gly
 245 250 255

Gly Glu Phe Gly Ser Trp Ala Phe Asn Leu Asn Pro Asp Gln Gly Glu
 260 265 270

Asn Asp Pro Gly Leu Trp Ile Ser Glu Pro Asn Gly Val Asp Ser Gly
 275 280 285

Leu Val Ala Ala Pro Val Met Asn Asn Gly Gly Asn Asp Ser Thr Ser
 290 295 300

Asn Ser Asp Ser Gln Pro Ile Ser Lys Leu Cys Asn Gly Ser Ser Val
 305 310 315 320

Glu Asn Pro Asn Pro Lys Val Leu Lys Ser Cys Glu Met Val Asn Phe
 325 330 335

Lys Asn Gly Ile Glu Asn Gly Gln Glu Glu Asp Ser Ser Asn Lys Lys
 340 345 350

Arg Ser Pro Val Ser Asn Asn Glu Glu Gly Met Leu Ser Phe Thr Ser
 355 360 365

Val Leu Pro Cys Asp Ser Asn His Ser Asp Leu Glu Ala Ser Val Ala
 370 375 380

Lys Glu Ala Glu Ser Asn Arg Val Val Val Glu Pro Glu Lys Lys Pro
 385 390 395 400

Arg Lys Arg Gly Arg Lys Pro Ala Asn Gly Arg Glu Glu Pro Leu Asn
 405 410 415

His Val Glu Ala Glu Arg Gln Arg Arg Glu Lys Leu Asn Gln Arg Phe
 420 425 430

Tyr Ser Leu Arg Ala Val Val Pro Asn Val Ser Lys Met Asp Lys Ala
 435 440 445

Ser Leu Leu Gly Asp Ala Ile Ser Tyr Ile Ser Glu Leu Lys Ser Lys
 450 455 460

Leu Gln Lys Ala Glu Ser Asp Lys Glu Glu Leu Gln Lys Gln Ile Asp
 465 470 475 480

Val Met Asn Lys Glu Ala Gly Asn Ala Lys Ser Ser Val Lys Asp Arg
 485 490 495

Lys Cys Leu Asn Gln Glu Ser Ser Val Leu Ile Glu Met Glu Val Asp
 500 505 510

Val Lys Ile Ile Gly Trp Asp Ala Met Ile Arg Ile Gln Cys Ser Lys
 515 520 525

MBI-17 Sequence Listing.ST25

Arg Asn His Pro Gly Ala Lys Phe Met Glu Ala Leu Lys Glu Leu Asp
530 535 540

Leu Glu Val Asn His Ala Ser Leu Ser Val Val Asn Asp Leu Met Ile
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Gln Gln Ala Thr Val Lys Met Gly Asn Gln Phe Phe Thr Gln Asp Gln
565 570 575

Leu Lys Val Ala Leu Thr Glu Lys Val Gly Glu Cys Pro
580 585

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Trp Thr Pro Glu Glu Asp Ile Ile Leu Val Ser Tyr Ile Gln Glu His
20 25 30

ggt cct gga aac tgg aga tct gtc cca aca cac aca ggt tta aga tgt 144
Gly Pro Gly Asn Trp Arg Ser Val Pro Thr His Thr Gly Leu Arg Cys
35 40 45

agc aag agc tgc aga ttg act aat tat ctt cga ccc ggt att 192
Ser Lys Ser Cys Arg Leu Arg Trp Thr Asn Tyr Leu Arg Pro Gly Ile
50 55 60

aag cgt gga aat ttt act gag cat gaa gag aag aca att gtt cat ctt 240
Lys Arg Gly Asn Phe Thr Glu His Glu Glu Lys Thr Ile Val His Leu
65 70 75 80

caa gcc ctt tta ggc aac aga tgg gca gcc ata gca tca tac ctt cca 288
Gln Ala Leu Leu Gly Asn Arg Trp Ala Ala Ile Ala Ser Tyr Leu Pro
85 90 95

gaa agg aca gac aat gat ata aag aac tat tgg aac act cac ttg aag 336
Glu Arg Thr Asp Asn Asp Ile Lys Asn Tyr Trp Asn Thr His Leu Lys
100 105 110

aag aag ctc aaa aag att aat gaa tct ggt gaa gaa gat aat gat ggt 384
Lys Lys Leu Lys Ile Asn Glu Ser Gly Glu Glu Asp Asn Asp Gly
115 120 125

gtc tct tca tca aac act agt tca caa aag aac cat caa agc act aac 432
Val Ser Ser Ser Asn Thr Ser Ser Gln Lys Asn His Gln Ser Thr Asn
130 135 140

aaa ggt caa tgg gaa aga aga ctt cag aca gac att aac atg gca aaa 480
Lys Gly Gln Trp Glu Arg Arg Leu Gln Thr Asp Ile Asn Met Ala Lys
145 150 155 160

caa gct ctt tgt gag gcc ttg tct tta gac aaa cca tca tcc act ctt 528
Gln Ala Leu Cys Glu Ala Leu Ser Leu Asp Lys Pro Ser Ser Thr Leu
165 170 175

tca tca tct tca tca tta ccg aca cca gta atc aca caa aac atc 576

MBI-17 Sequence Listing ST25

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180	185	190
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Arg Asn Phe Ser Ser Ala Leu Leu Asp Arg Cys Tyr Asp Pro Ser Ser		
195	200	205
tct tct tca tct acc aca acc acc act aca agc aac act act aat cca		672
Ser Ser Ser Thr Thr Thr Thr Ser Asn Thr Thr Asn Pro		
210	215	220
tac cca tca ggg gta tat gcg tca agt gct gag aac atc gcc cgg ttg		720
Tyr Pro Ser Gly Val Tyr Ala Ser Ser Ala Glu Asn Ile Ala Arg Leu		
225	230	235
240		
ctt caa gat ttc atg aaa gac aca ccc aag gct tta act tta tca tct		768
Leu Gln Asp Phe Met Lys Asp Thr Pro Lys Ala Leu Thr Leu Ser Ser		
245	250	255
tca tct ccg gtt tca gag act gga cca ctc act gct gca gtc tcg gaa		816
Ser Ser Pro Val Ser Glu Thr Gly Pro Leu Thr Ala Ala Val Ser Glu		
260	265	270
gaa ggt gga gaa ggg ttt gaa caa tct ttc ttc agc ttc aat tca atg		864
Glu Gly Gly Glu Gly Phe Glu Gln Ser Phe Phe Ser Phe Asn Ser Met		
275	280	285
gac gaa act caa aac ttg act cag gag aca agc ttc ttc cat gat caa		912
Asp Glu Thr Gln Asn Leu Thr Gln Glu Thr Ser Phe Phe His Asp Gln		
290	295	300
gtg atc aaa ccg gaa ata aca atg gac caa gat cat ggt cta ata tca		960
Val Ile Lys Pro Glu Ile Thr Met Asp Gln Asp His Gly Leu Ile Ser		
305	310	315
320		
caa ggg tct ctg tct ttg ttt gag aaa tgg tta ttt gat gag caa agc		1008
Gln Gly Ser Leu Ser Leu Phe Glu Lys Trp Leu Phe Asp Glu Gln Ser		
325	330	335
cac gag atg gtt ggt atg gca cta gca gga caa gaa ggg atg ttc tag		1056
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Gly Pro Gly Asn Trp Arg Ser Val Pro Thr His Thr Gly Leu Arg Cys		
35	40	45
Ser Lys Ser Cys Arg Leu Arg Trp Thr Asn Tyr Leu Arg Pro Gly Ile		
50	55	60
Lys Arg Gly Asn Phe Thr Glu His Glu Glu Lys Thr Ile Val His Leu		
65	70	75
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Gln Ala Leu Leu Gly Asn Arg Trp Ala Ala Ile Ala Ser Tyr Leu Pro		
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MBI-17 Sequence Listing, ST25

Glu Arg Thr Asp Asn Asp Ile Lys Asn Tyr Trp Asn Thr His Leu Lys
100 105 110

Lys Lys Leu Lys Lys Ile Asn Glu Ser Gly Glu Glu Asp Asn Asp Gly
115 120 125

Val Ser Ser Ser Asn Thr Ser Ser Gln Lys Asn His Gln Ser Thr Asn
130 135 140

Lys Gly Gln Trp Glu Arg Arg Leu Gln Thr Asp Ile Asn Met Ala Lys
145 150 155 160

Gln Ala Leu Cys Glu Ala Leu Ser Leu Asp Lys Pro Ser Ser Thr Leu
165 170 175

Ser Ser Ser Ser Ser Leu Pro Thr Pro Val Ile Thr Gln Gln Asn Ile
180 185 190

Arg Asn Phe Ser Ser Ala Leu Leu Asp Arg Cys Tyr Asp Pro Ser Ser
195 200 205

Ser Ser Ser Ser Thr Thr Thr Thr Thr Ser Asn Thr Thr Asn Pro
210 215 220

Tyr Pro Ser Gly Val Tyr Ala Ser Ser Ala Glu Asn Ile Ala Arg Leu
225 . . . 230 . . . 235 . . . 240

Leu Gln Asp Phe Met Lys Asp Thr Pro Lys Ala Leu Thr Leu Ser Ser
245 250 255

Ser Ser Pro Val Ser Glu Thr Gly Pro Leu Thr Ala Ala Val Ser Glu
260 265 270

Glu Gly Gly Glu Gly Phe Glu Gln Ser Phe Phe Ser Phe Asn Ser Met
275 280 285

Asp Glu Thr Gln Asn Leu Thr Gln Glu Thr Ser Phe Phe His Asp Gln
290 295 300

Val Ile Lys Pro Glu Ile Thr Met Asp Gln Asp His Gly Leu Ile Ser
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Gln Gly Ser Leu Ser Leu Phe Glu Lys Trp Leu Phe Asp Glu Gln Ser
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His Glu Met Val Gly Met Ala Leu Ala Gly Gln Glu Gly Met Phe

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MBI-17 Sequence Listing.ST25

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 Met Asp Thr Asn Thr Ser
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 gga gaa gaa tta tta gct aag gca aga aag cca tat aca ata aca aag 403
 Gly Glu Glu Leu Leu Ala Lys Ala Arg Lys Pro Tyr Thr Ile Thr Lys
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 cag cga gag cga tgg act gag gat gag cat gag agg ttt cta gaa gcc 451
 Gln Arg Glu Arg Trp Thr Glu Asp Glu His Glu Arg Phe Leu Glu Ala
 25 30 35
 ttg agg ctt tat gga aga gct tgg caa cga att gaa gaa cat att ggg 499
 Leu Arg Leu Tyr Gly Arg Ala Trp Gln Arg Ile Glu Glu His Ile Gly
 40 45 50
 aca aag act gct gtt cag atc aga agt cat gca caa aag ttc ttc aca 547
 Thr Lys Thr Ala Val Gln Ile Arg Ser His Ala Gln Lys Phe Phe Thr
 55 60 65 70
 aag ttg gag aaa gag gct gaa gtt aaa ggc atc cct gtt tgc caa gct 595
 Lys Leu Glu Lys Glu Ala Glu Val Lys Gly Ile Pro Val Cys Gln Ala
 75 80 85
 ttg gac ata gaa att ccg cct cct cgt cct aaa cga aaa ccc aat act 643
 Leu Asp Ile Glu Ile Pro Pro Arg Pro Lys Arg Lys Pro Asn Thr
 90 95 100
 cct tat cct cga aaa cct ggg aac aac ggt aca tct tcc tct caa gta 691
 Pro Tyr Pro Arg Lys Pro Gly Asn Asn Gly Thr Ser Ser Ser Gln Val
 105 110 115
 tca tca gca aaa gat gca aaa ctt gtt tca tcg gcc tct tct tca cag 739
 Ser Ser Ala Lys Asp Ala Lys Leu Val Ser Ser Ala Ser Ser Gln
 120 125 130
 ttg aat cag gcg ttc ttg gat ttg gaa aaa atg ccg ttc tct gag aaa 787
 Leu Asn Gln Ala Phe Leu Asp Leu Glu Lys Met Pro Phe Ser Glu Lys
 135 140 145 150
 aca tca act gga aaa gaa aat caa gat gag aat tgc tcg ggt gtt tct 835
 Thr Ser Thr Gly Lys Glu Asn Gln Asp Glu Asn Cys Ser Gly Val Ser
 155 160 165
 act gtg aac aag tat ccc tta cca acg aaa cag gta agt ggc gac att 883
 Thr Val Asn Lys Tyr Pro Leu Pro Thr Lys Gln Val Ser Gly Asp Ile
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 gaa aca agt aag acc tca act gtg gac aac gcg gtt caa gat gtt ccc 931
 Glu Thr Ser Lys Thr Ser Thr Val Asp Asn Ala Val Gln Asp Val Pro
 185 190 195
 aag aag aac aaa gac aaa gat ggt aac gat ggt act act gtg cac agc 979
 Lys Lys Asn Lys Asp Lys Asp Gly Asn Asp Gly Thr Thr Val His Ser
 200 205 210
 atg caa aac tac cct tgg cat ttc cac gca gat att gtg aac ggg aat 1027
 Met Gln Asn Tyr Pro Trp His Phe His Ala Asp Ile Val Asn Gly Asn
 215 220 225 230
 ata gca aaa tgc cct caa aat cat ccc tca ggt atg gta tct caa gac 1075
 Ile Ala Lys Cys Pro Gln Asn His Pro Ser Gly Met Val Ser Gln Asp
 235 240 245

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caa gct aca aca gca tct gct act act aca gct tct cat caa gcg ttt Gln Ala Thr Thr Ala Ser Ala Thr Thr Ala Ser His Gln Ala Phe 265 270 275	1171
cca gct tgt cat tca cag gat gat tac cgt tcg ttt ctc cag ata tca Pro Ala Cys His Ser Gln Asp Asp Tyr Arg Ser Phe Leu Gln Ile Ser 280 285 290	1219
tct act ttc tcc aat ctt att atg tca act ctc cta cag aat cct gca Ser Thr Phe Ser Asn Leu Ile Met Ser Thr Leu Leu Gln Asn Pro Ala 295 300 305 310	1267
gct cat gct gca gct aca ttc gct gct tcg gtc tgg cct tat gcg agt Ala His Ala Ala Ala Thr Phe Ala Ala Ser Val Trp Pro Tyr Ala Ser 315 320 325	1315
gtc ggg aat tct ggt gat tca tca acc cca atg agc tct tct cct cca Val Gly Asn Ser Gly Asp Ser Ser Thr Pro Met Ser Ser Pro Pro 330 335 340	1363
agt ata act gcc att gcc gct gct aca gta gct gct gca act gct tgg Ser Ile Thr Ala Ile Ala Ala Ala Thr Val Ala Ala Ala Thr Ala Trp 345 350 355	1411
tgg gct tct cat gga ctt ctt cct gta tgc gct cca gct cca ata aca Trp Ala Ser His Gly Leu Leu Pro Val Cys Ala Pro Ala Pro Ile Thr 360 365 370	1459
tgt gtt cca ttc tca act gtt gca gtt cca act cca gca atg act gaa Cys Val Pro Phe Ser Thr Val Ala Val Pro Thr Pro Ala Met Thr Glu 375 380 385 390	1507
atg gat acc gtt gaa aat act caa ccg ttt gag aaa caa aac aca gct Met Asp Thr Val Glu Asn Thr Gln Pro Phe Glu Lys Gln Asn Thr Ala 395 400 405	1555
ctg caa gat caa acc ttg gct tcg aaa tct cca gct tca tca tct gat Leu Gln Asp Gln Thr Leu Ala Ser Lys Ser Pro Ala Ser Ser Ser Asp 410 415 420	1603
gat tca gat gag act gga gta acc aag cta aat gcc gac tca aaa acc Asp Ser Asp Glu Thr Gly Val Thr Lys Leu Asn Ala Asp Ser Lys Thr 425 430 435	1651
aat gat gat aaa att gag gag gtt gtt act gcc gct gtg cat gac Asn Asp Asp Lys Ile Glu Glu Val Val Val Thr Ala Ala Val His Asp 440 445 450	1699
tca aac act gcc cag aag aaa aat ctt gtg gac cgc tca tcg tgt ggc Ser Asn Thr Ala Gln Lys Lys Asn Leu Val Asp Arg Ser Ser Cys Gly 455 460 465 470	1747
tca aat aca cct tca ggg agt gac gca gaa act gat gca tta gat aaa Ser Asn Thr Pro Ser Gly Ser Asp Ala Glu Thr Asp Ala Leu Asp Lys 475 480 485	1795
atg gag aaa gat aaa gag gat gtg aag gag aca gat gag aat gag cca Met Glu Lys Asp Lys Glu Asp Val Lys Glu Thr Asp Glu Asn Gln Pro 490 495 500	1843
gat gtt att gag tta aat aac cgt aag att aaa atg aga gac aac aac Asp Val Ile Glu Leu Asn Asn Arg Lys Ile Lys Met Arg Asp Asn Asn 505 510 515	1891
agc aac aac aat gca act act gat tcg tgg aag gaa gtc tcc gaa gag Ser Asn Asn Asn Ala Thr Thr Asp Ser Trp Lys Glu Val Ser Glu Glu 520 525 530	1939
ggc cgt ata gcg ttt cag gct ctc ttt gca aga gaa gca aga ttg cct caa Gly Arg Ile Ala Phe Gln Ala Leu Phe Ala Arg Glu Arg Leu Pro Gln 535 540 545 550	1987

MBI-17 Sequence Listing.ST25

agc ttt tcg cct cct caa gtg gca gag aat gtg aat aga aaa caa agt Ser Phe Ser Pro Pro Gln Val Ala Glu Asn Val Asn Arg Lys Gln Ser 555 560 565	2035
gac acg tca atg cca ttg gct cct aat ttc aaa agc cag gat tct tgt Asp Thr Ser Met Pro Leu Ala Pro Asn Phe Lys Ser Gln Asp Ser Cys 570 575 580	2083
gct gca gac caa gaa gga gta gta atg atc ggt gtt gga aca tgc aag Ala Ala Asp Gln Glu Gly Val Val Met Ile Gly Val Gly Thr Cys Lys 585 590 595	2131
agt ctt aaa acg aga cag aca gga ttt aag cca tac aag aga tgt tca Ser Ieu Lys Thr Arg Gln Thr Gly Phe Lys Pro Tyr Lys Arg Cys Ser 600 605 610	2179
atg gaa gtg aaa gag agc caa gtt ggg aac ata aac aat caa agt gat Met Glu Val Lys Glu Ser Gln Val Gly Asn Ile Asn Asn Gln Ser Asp 615 620 625 630	2227
gaa aaa gtc tgc aaa agg ctt cga ttg gaa gga gaa gct tct aca tga Glu Lys Val Cys Lys Arg Leu Arg Leu Glu Gly Glu Ala Ser Thr 635 640 645	2275
cagacttgaa ggtaaaaaaaaa aaacatccac atttttatca atatctttaa atcttagtgg atctttgc ttctccaatc tttatgaaag agacttttaa ttttccttcc gaacatttct tttgtcatgt caggttctgt accatattac cccatgtctt gtctttgtc tctgtttgtg tatgctactt gtggctata tgtcatctgc tactactgtt aattaaccat taagcaatgg atttgtcttt a	2335 2395 2455 2515 2526
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Met Asp Thr Asn Thr Ser Gly Glu Glu Leu Leu Ala Lys Ala Arg Lys 1 5 10 15	
Pro Tyr Thr Ile Thr Lys Gln Arg Glu Arg Trp Thr Glu Asp Glu His 20 25 30	
Glu Arg Phe Leu Glu Ala Leu Arg Leu Tyr Gly Arg Ala Trp Gln Arg 35 40 45	
Ile Glu Glu His Ile Gly Thr Lys Thr Ala Val Gln Ile Arg Ser His 50 55 60	
Ala Gln Lys Phe Phe Thr Lys Leu Glu Lys Glu Ala Glu Val Lys Gly 65 70 75 80	
Ile Pro Val Cys Gln Ala Leu Asp Ile Glu Ile Pro Pro Pro Arg Pro 85 90 95	
Lys Arg Lys Pro Asn Thr Pro Tyr Pro Arg Lys Pro Gly Asn Asn Gly 100 105 110	
Thr Ser Ser Ser Gln Val Ser Ser Ala Lys Asp Ala Lys Leu Val Ser 115 120 125	

MBI-17 Sequence Listing ST25

Ser Ala Ser Ser Ser Gln Leu Asn Gln Ala Phe Leu Asp Leu Glu Lys
130 135 140

Met Pro Phe Ser Glu Lys Thr Ser Thr Gly Lys Glu Asn Gln Asp Glu
145 150 155 160

Asn Cys Ser Gly Val Ser Thr Val Asn Lys Tyr Pro Leu Pro Thr Lys
165 170 175

Gln Val Ser Gly Asp Ile Glu Thr Ser Lys Thr Ser Thr Val Asp Asn
180 185 190

Ala Val Gln Asp Val Pro Lys Lys Asn Lys Asp Lys Asp Gly Asn Asp
195 200 205

Gly Thr Thr Val His Ser Met Gln Asn Tyr Pro Trp His Phe His Ala
210 215 220

Asp Ile Val Asn Gly Asn Ile Ala Lys Cys Pro Gln Asn His Pro Ser
225 230 235 240

Gly Met Val Ser Gln Asp Phe Met Phe His Pro Met Arg Glu Glu Thr
245 250 255

His Gly His Ala Asn Leu Gln Ala Thr Thr Ala Ser Ala Thr Thr Thr
260 265 270

Ala Ser His Gln Ala Phe Pro Ala Cys His Ser Gln Asp Asp Tyr Arg
275 280 285

Ser Phe Leu Gln Ile Ser Ser Thr Phe Ser Asn Leu Ile Met Ser Thr
290 295 300

Leu Leu Gln Asn Pro Ala Ala His Ala Ala Ala Thr Phe Ala Ala Ser
305 310 315 320

Val Trp Pro Tyr Ala Ser Val Gly Asn Ser Gly Asp Ser Ser Thr Pro
325 330 335

Met Ser Ser Ser Pro Pro Ser Ile Thr Ala Ile Ala Ala Ala Thr Val
340 345 350

Ala Ala Ala Thr Ala Trp Trp Ala Ser His Gly Leu Leu Pro Val Cys
355 360 365

Ala Pro Ala Pro Ile Thr Cys Val Pro Phe Ser Thr Val Ala Val Pro
370 375 380

Thr Pro Ala Met Thr Glu Met Asp Thr Val Glu Asn Thr Gln Pro Phe
385 390 395 400

Glu Lys Gln Asn Thr Ala Leu Gln Asp Gln Thr Leu Ala Ser Lys Ser
405 410 415

Pro Ala Ser Ser Ser Asp Asp Ser Asp Glu Thr Gly Val Thr Lys Leu
420 425 430

MBI-17 Sequence Listing.ST25

Asn Ala Asp Ser Lys Thr Asn Asp Asp Lys Ile Glu Glu Val Val Val
 435 440 445

Thr Ala Ala Val His Asp Ser Asn Thr Ala Gln Lys Lys Asn Leu Val
 450 455 460

Asp Arg Ser Ser Cys Gly Ser Asn Thr Pro Ser Gly Ser Asp Ala Glu
 465 470 475 480

Thr Asp Ala Leu Asp Lys Met Glu Lys Asp Lys Glu Asp Val Lys Glu
 485 490 495

Thr Asp Glu Asn Gln Pro Asp Val Ile Glu Leu Asn Asn Arg Lys Ile
 500 505 510

Lys Met Arg Asp Asn Asn Ser Asn Asn Ala Thr Thr Asp Ser Trp
 515 520 525

Lys Glu Val Ser Glu Glu Gly Arg Ile Ala Phe Gln Ala Leu Phe Ala
 530 535 540

Arg Glu Arg Leu Pro Gln Ser Phe Ser Pro Pro Gln Val Ala Glu Asn
 545 550 555 560

Val Asn Arg Lys Gln Ser Asp Thr Ser Met Pro Leu Ala Pro Asn Phe
 565 570 575

Lys Ser Gln Asp Ser Cys Ala Ala Asp Gln Glu Gly Val Val Met Ile
 580 585 590

Gly Val Gly Thr Cys Lys Ser Leu Lys Thr Arg Gln Thr Gly Phe Lys
 595 600 605

Pro Tyr Lys Arg Cys Ser Met Glu Val Lys Glu Ser Gln Val Gly Asn.
 610 615 620

Ile Asn Asn Gln Ser Asp Glu Lys Val Cys Lys Arg Leu Arg Leu Glu
 625 630 635 640

Gly Glu Ala Ser Thr
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 <222> (1)..(228)
 <223> G682

<400> 33
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 Met Asp Asn His Arg Arg Thr Lys Gln Pro Lys Thr Asn Ser Ile Val
 1 5 10 15

act tct tct gaa gaa gtg agt agt ctt gag tgg gaa gtt gtg aac
 Thr Ser Ser Glu Glu Val Ser Ser Leu Glu Trp Glu Val Val Asn

48

96

MBI-17 Sequence Listing.ST25
25 30

20

25

30

atg agt caa gaa gaa gaa gat ttg gtc tct cga atg cat aag ctt gtc 144
 Met Ser Gln Glu Glu Glu Asp Leu Val Ser Arg Met His Lys Leu Val
 35 40 45

ggt gac agg tgg gaa ctg ata gct ggg agg atc cca gga aga acc gct 192
 Gly Asp Arg Trp Glu Leu Ile Ala Gly Arg Ile Pro Gly Arg Thr Ala
 50 55 60

<210> 34
<211> 75
<212> PRT
<213> *Arabidopsis thaliana*

<400> 34

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Thr Ser Ser Ser Glu Glu Val Ser Ser Leu Glu Trp Glu Val Val Asn
20 25 30

Met Ser Gln Glu Glu Glu Asp Leu Val Ser Arg Met His Lys Leu Val
35 40 45

Gly Asp Arg Trp Glu Leu Ile Ala Gly Arg Ile Pro Gly Arg Thr Ala
50 55 60

Gly Glu Ile Glu Arg Phe Trp Val Met Lys Asn
65 70 75

<210> 35
<211> 584
<212> DNA
<213> *Arabidopsis thaliana*

<220>
<221> CDS
<222> (157)..(441)
<223> G225

gcg gaa aaa atg gat aaa cga cga cg_g aga cag agc aaa gcc aag gct 222
 Ala Glu Lys Met Asp Lys Arg Arg Arg Gln Ser Lys Ala Lys Ala
 10 15 20

tct tgt tcc gaa gag gtg agt agt atc gaa tgg gaa gct gtg aag atg
 Ser Cys Ser Glu Glu Val Ser Ser Ile Glu Trp Glu Ala Val Lys Met
 25 30 35

tca gaa gaa gaa gaa gat ctc att tct cg^g atg tat aaa ctc gtt ggc 318
 Ser Glu Glu Glu Glu Asp Leu Ile Ser Arg Met Tyr Lys Leu Val Gly
 40 45 50

gac agg tgg gag ttg atc gcc gga agg atc ccg gga cgg acg ccg gag 366

MBI-17 Sequence Listing ST25

Asp Arg Trp Glu Leu Ile Ala Gly Arg Ile Pro Gly Arg Thr Pro Glu
 55 60 65 70

gag ata gag aga tat tgg ctt atg aaa cac ggc gtc gtt ttt gcc aac 414
 Glu Ile Glu Arg Tyr Trp Leu Met Lys His Gly Val Val Phe Ala Asn
 75 80 85

aga cga aga gac ttt ttt agg aaa tga tttttttgt ttggattaaa 461
 Arg Arg Arg Asp Phe Phe Arg Lys
 90

agaaaaatttt cctctcctta attcacaaga caagaaaaaa agggaaatgta cctgtccttg 521
 aattactatt ttggaatgta taattatcta tatataaag aagaaaaat tgcttagaa 581
 ttt 584

<210> 36
<211> 94
<212> PRT
<213> Arabidopsis thaliana

<400> 36

Met Phe Arg Ser Asp Lys Ala Glu Lys Met Asp Lys Arg Arg Arg Arg
 1 5 10 15

Gln Ser Lys Ala Lys Ala Ser Cys Ser Glu Glu Val Ser Ser Ile Glu
 20 25 30

Trp Glu Ala Val Lys Met Ser Glu Glu Glu Glu Asp Leu Ile Ser Arg
 35 40 45

Met Tyr Lys Leu Val Gly Asp Arg Trp Glu Leu Ile Ala Gly Arg Ile
 50 55 60

Pro Gly Arg Thr Pro Glu Glu Ile Glu Arg Tyr Trp Leu Met Lys His
 65 70 75 80

Gly Val Val Phe Ala Asn Arg Arg Asp Phe Phe Arg Lys
 85 90

<210> 37
<211> 1369
<212> DNA
<213> Arabidopsis thaliana

<220>
<221> CDS
<222> (104)..(1174)
<223> G678

<400> 37
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aacaagtgaa gtgagtgatt ttataataatc gccggccgag aga atg gga agg gcg 115
Met Gly Arg Ala
1

ccg tgc tgt gag aag gta gga att aag aag ggg cgc tgg acg gcg gag 163
Pro Cys Cys Glu Lys Val Gly Ile Lys Lys Gly Arg Trp Thr Ala Glu
5 10 15 20

gaa gac cgg act ctc tcc gac tac att cag tcc aac ggc gaa gga tca 211
Glu Asp Arg Thr Leu Ser Asp Tyr Ile Gln Ser Asn Gly Glu Gly Ser
25 30 35

MBI-17 Sequence Listing, ST25

tgg cgt tct ctt ccc aaa aat gcc ggg cta aag aga tgt gga aag agc Trp Arg Ser Leu Pro Lys Asn Ala Gly Leu Lys Arg Cys Gly Lys Ser 40 45 50	259
tgt aga ttg aga tgg ata aac tat ttg aga tca gac atc aag aga gga Cys Arg Leu Arg Trp Ile Asn Tyr Leu Arg Ser Asp Ile Lys Arg Gly 55 60 65	307
aac ata act ccc gaa gaa gag gac gtc att gtt aaa ctg cat tcc act Asn Ile Thr Pro Glu Glu Asp Val Ile Val Lys Leu His Ser Thr 70 75 80	355
ttg gga acc agg tgg tca aca att gcg agc aat cta ccg gga aga aca Leu Gly Thr Arg Trp Ser Thr Ile Ala Ser Asn Leu Pro Gly Arg Thr 85 90 95 100	403
gac aac gaa ata aaa aac tat tgg aat tct cat ctc agc cgt aaa ctc Asp Asn Glu Ile Lys Asn Tyr Trp Asn Ser His Leu Ser Arg Lys Leu 105 110 115	451
cac ggt tac ttc aga aaa cca act gtc gcc aat acc gtc gag aat gcg His Gly Tyr Phe Arg Lys Pro Thr Val Ala Asn Thr Val Glu Asn Ala 120 125 130	499
cct ccg cct aag cgt aga cct gga aga acc agc aga tcc gcc atg Pro Pro Pro Pro Lys Arg Arg Pro Gly Arg Thr Ser Arg Ser Ala Met 135 140 145	547
aaa ccc aaa ttt atc cta aac cct aaa aac cac aaa acc cct aat tct Lys Pro Lys Phe Ile Leu Asn Pro Lys Asn His Lys Thr Pro Asn Ser 150 155 160	595
ttt aaa gca aac aaa agt gac atc gtt ttg cca act acg aca ata gag Phe Lys Ala Asn Lys Ser Asp Ile Val Leu Pro Thr Thr Thr Ile Glu 165 170 175 180	643
aat gga gag gga gac aaa gaa gac gca tta atg gtg ttg tca agt agt Asn Gly Glu Gly Asp Lys Glu Asp Ala Leu Met Val Leu Ser Ser Ser 185 190 195	691
agc tta agt gga gca gag gaa ccc ggt tta gga cca tgt ggt tat gga Ser Leu Ser Gly Ala Glu Glu Pro Gly Leu Gly Pro Cys Gly Tyr Gly 200 205 210	739
gac gat ggc gat tgt aac cca agc att aat ggc gac gat gga gct ttg Asp Asp Gly Asp Cys Asn Pro Ser Ile Asn Gly Asp Asp Gly Ala Leu 215 220 225	787
tgt ctc aat gac gac att ttc gat tct tgt ttt cta ttg gac gac tct Cys Leu Asn Asp Asp Ile Phe Asp Ser Cys Phe Leu Leu Asp Asp Ser 230 235 240	835
cat gct gtc cac gtg tcc tca tgt gag tcg aac aac gta aaa aac tct His Ala Val His Val Ser Ser Cys Glu Ser Asn Asn Val Lys Asn Ser 245 250 255 260	883
gag cca tat gga ggg atg tca gtt ggg cac aaa aat atc gaa acg atg Glu Pro Tyr Gly Gly Met Ser Val Gly His Lys Asn Ile Glu Thr Met 265 270 275	931
gct gat gat ttc gtt gac tgg gac ttt gta tgg aga gaa ggt caa acc Ala Asp Asp Phe Val Asp Trp Asp Phe Val Trp Arg Glu Gly Gln Thr 280 285 290	979
ctt tgg gac gaa aaa gag gat ctt gat tcg gtt ttg tcg agg ctg tta Leu Trp Asp Glu Lys Glu Asp Leu Asp Ser Val Leu Ser Arg Leu Leu 295 300 305	1027
gat gga gag gaa atg gaa tct gag atc aga caa agg gac tcc aac gac Asp Gly Glu Glu Met Glu Ser Glu Ile Arg Gln Arg Asp Ser Asn Asp 310 315 320	1075
ttt gga gaa ccg ttg gat att gac gaa gaa aac aag atg gct gct tgg Phe Gly Glu Pro Leu Asp Ile Asp Glu Glu Asn Lys Met Ala Ala Trp 325 330 335	1123

MBI-17 Sequence Listing ST25

ctt ttt tcc tta aaa att tta ccc cct tcc ttt tcc ctt ttc ccc ctt
 Leu Phe Ser Leu Lys Ile Leu Pro Pro Ser Phe Ser Leu Phe Pro Leu
 345 350 355

taa tttttaccaa aaccggccctt tgccagatcc tgtccgtttt tccattaaac 1224

ctttttctcc ccctacccctc ctttttttat tttaattttt ttttttttcc ttttttttcc 1284

ctttccctttt ttaattccga tttttggcggttgccaattt aaccaaaatata aatccatcc 1344

taaaaaaaaaaa aaaaaaaaaaaa aaaaaa 1369

<210> 38

<211> 356

<212> PRT

<213> Arabidopsis thaliana

<400> 38

Met Gly Arg Ala Pro Cys Cys Glu Lys Val Gly Ile Lys Lys Gly Arg
 1 5 10 15

Trp Thr Ala Glu Glu Asp Arg Thr Leu Ser Asp Tyr Ile Gln Ser Asn
 20 25 30

Gly Glu Gly Ser Trp Arg Ser Leu Pro Lys Asn Ala Gly Leu Lys Arg
 35 40 45

Cys Gly Lys Ser Cys Arg Leu Arg Trp Ile Asn Tyr Leu Arg Ser Asp
 50 55 60

Ile Lys Arg Gly Asn Ile Thr Pro Glu Glu Asp Val Ile Val Lys
 65 70 75 80

Leu His Ser Thr Leu Gly Thr Arg Trp Ser Thr Ile Ala Ser Asn Leu
 85 90 95

Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Tyr Trp Asn Ser His Leu
 100 105 110

Ser Arg Lys Leu His Gly Tyr Phe Arg Lys Pro Thr Val Ala Asn Thr
 115 120 125

Val Glu Asn Ala Pro Pro Pro Lys Arg Arg Pro Gly Arg Thr Ser
 130 135 140

Arg Ser Ala Met Lys Pro Lys Phe Ile Leu Asn Pro Lys Asn His Lys
 145 150 155 160

Thr Pro Asn Ser Phe Lys Ala Asn Lys Ser Asp Ile Val Leu Pro Thr
 165 170 175

Thr Thr Ile Glu Asn Gly Glu Asp Lys Glu Asp Ala Leu Met Val
 180 185 190

Leu Ser Ser Ser Ser Leu Ser Gly Ala Glu Glu Pro Gly Leu Gly Pro
 195 200 205

Cys Gly Tyr Gly Asp Asp Gly Asp Cys Asn Pro Ser Ile Asn Gly Asp
 210 215 220

MBI-17 Sequence Listing ST25

Asp Gly Ala Leu Cys Leu Asn Asp Asp Ile Phe Asp Ser Cys Phe Leu
 225 230 235 240

Leu Asp Asp Ser His Ala Val His Val Ser Ser Cys Glu Ser Asn Asn
 245 250 255

Val Lys Asn Ser Glu Pro Tyr Gly Gly Met Ser Val Gly His Lys Asn
 260 265 270

Ile Glu Thr Met Ala Asp Asp Phe Val Asp Trp Asp Phe Val Trp Arg
 275 280 285

Glu Gly Gln Thr Leu Trp Asp Glu Lys Glu Asp Leu Asp Ser Val Leu
 290 295 300

Ser Arg Leu Leu Asp Gly Glu Glu Met Glu Ser Glu Ile Arg Gln Arg
 305 310 315 320

Asp Ser Asn Asp Phe Gly Glu Pro Leu Asp Ile Asp Glu Glu Asn Lys
 325 330 335

Met Ala Ala Trp Leu Phe Ser Leu Lys Ile Leu Pro Pro Ser Phe Ser
 340 345 350

Leu Phe Pro Leu
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<210> 39
 <211> 1046
 <212> DNA
 <213> Arabidopsis thaliana

<220>
 <221> CDS
 <222> (46)..(867)
 <223> G233

<400> 39
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 Met Gly Arg Ala
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cca tgc tgt gag aag atg ggg ttg aag aga gga cca tgg aca cct gaa 105
 Pro Cys Cys Glu Lys Met Gly Leu Lys Arg Gly Pro Trp Thr Pro Glu
 5 10 15 20

gaa gat caa atc ttg gtc tct ttt atc ctc aac cat gga cat agt aac 153
 Glu Asp Gln Ile Leu Val Ser Phe Ile Leu Asn His Gly His Ser Asn
 25 30 35

tgg cga gcc ctc cct aag caa gct ggt ctt ttg aga tgt gga aaa agc 201
 Trp Arg Ala Leu Pro Lys Gln Ala Gly Leu Leu Arg Cys Gly Lys Ser
 40 45 50

tgt aga ctt agg tgg atg aac tat tta aag cct gat att aaa cgt ggc 249
 Cys Arg Leu Arg Trp Met Asn Tyr Leu Lys Pro Asp Ile Lys Arg Gly
 55 60 65

aat ttc acc aaa gaa gag gaa gat gct atc atc agc tta cac caa ata 297
 Asn Phe Thr Lys Glu Glu Asp Ala Ile Ile Ser Leu His Gln Ile
 70 75 80

ctt ggc aat aga tgg tca gcg att gca gca aaa ctg cct gga aga acc 345

MBI-17 Sequence Listing ST25

Leu Gly Asn Arg Trp Ser Ala Ile Ala Ala Lys Leu Pro Gly Arg Thr
85 90 95 100

gat aac gag atc aag aac gta tgg cac act cac ttg aag aag aga ctc 393
Asp Asn Glu Ile Lys Asn Val Trp His Thr His Leu Lys Lys Arg Leu
105 110 115

gaa gat tat caa cca gct aaa cct aag acc agc aac aaa aag aag ggt 441
Glu Asp Tyr Gln Pro Ala Lys Pro Lys Thr Ser Asn Lys Lys Lys Gly
120 125 130

act aaa cca aaa tct gaa tcc gta ata acg agc tcg aac agt act aga 489
Thr Lys Pro Lys Ser Glu Ser Val Ile Thr Ser Ser Asn Ser Thr Arg
135 140 145

agc gaa tcg gag cta gca gat tca tca aac cct tct gga gaa agc tta 537
Ser Glu Ser Glu Leu Ala Asp Ser Ser Asn Pro Ser Gly Glu Ser Leu
150 155 160

ttt tcg aca tcg cct tcg aca agt gag gtt tct tcg atg aca ctc ata 585
Phe Ser Thr Ser Pro Ser Thr Ser Glu Val Ser Ser Met Thr Leu Ile
165 170 175 180

agc cac gac ggc tat agc aac gag att aat atg gat aac aaa ccg gga 633
Ser His Asp Gly Tyr Ser Asn Glu Ile Asn Met Asp Asn Lys Pro Gly
185 190 195

gat atc agt act atc gat caa gaa tgt gtt tct ttc gaa act ttt ggt 681
Asp Ile Ser Thr Ile Asp Gln Glu Cys Val Ser Phe Glu Thr Phe Gly
200 205 210

gcg gat atc gat gaa agc ttc tgg aaa gag aca ctg tat agc caa gat 729
Ala Asp Ile Asp Glu Ser Phe Trp Lys Glu Thr Leu Tyr Ser Gln Asp
215 220 225

gaa cac aac tac gta tcg aat gac cta gaa gtc gct ggt tta gtt gag 777
Glu His Asn Tyr Val Ser Asn Asp Leu Glu Val Ala Gly Leu Val Glu
230 235 240

ata caa caa gag ttt caa aac ttg ggc tcc gct aat aat gag atg att 825
Ile Gln Gln Glu Phe Gln Asn Leu Gly Ser Ala Asn Asn Glu Met Ile
245 250 255 260

ttt gac agt gag atg gaa ctt ctg gtt cga tgt att ggc tag 867
Phe Asp Ser Glu Met Glu Leu Leu Val Arg Cys Ile Gly
265 270

aaccggcggg gaacaagatc tcttagccgg gctctagtttta acatgtttga ggagtaaagt 927

gaaatggtgc aaatttagtttta aggctaagaa attcaaaagc ttttgtttac cgagaaaaaa 987

acacactcta actcttgatg tgatgttagtt agtgttattaa tttagaggctg cgttttcaa 1046

<210> 40
<211> 273
<212> PRT
<213> Arabidopsis thaliana

<400> 40

Met Gly Arg Ala Pro Cys Cys Glu Lys Met Gly Leu Lys Arg Gly Pro
1 5 10 15

Trp Thr Pro Glu Glu Asp Gln Ile Leu Val Ser Phe Ile Leu Asn His
20 25 30

Gly His Ser Asn Trp Arg Ala Leu Pro Lys Gln Ala Gly Leu Leu Arg
35 40 45

Cys Gly Lys Ser Cys Arg Leu Arg Trp Met Asn Tyr Leu Lys Pro Asp
50 55 60

MBI-17 Sequence Listing.ST25

Ile Lys Arg Gly Asn Phe Thr Lys Glu Glu Glu Asp Ala Ile Ile Ser
 65 70 75 80

Leu His Gln Ile Leu Gly Asn Arg Trp Ser Ala Ile Ala Ala Lys Leu
 85 90 95

Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Val Trp His Thr His Leu
 100 105 110

Lys Lys Arg Leu Glu Asp Tyr Gln Pro Ala Lys Pro Lys Thr Ser Asn
 115 120 125

Lys Lys Lys Gly Thr Lys Pro Lys Ser Glu Ser Val Ile Thr Ser Ser
 130 135 140

Asn Ser Thr Arg Ser Glu Ser Glu Leu Ala Asp Ser Ser Asn Pro Ser
 145 150 155 160

Gly Glu Ser Leu Phe Ser Thr Ser Pro Ser Thr Ser Glu Val Ser Ser
 165 170 175

Met Thr Leu Ile Ser His Asp Gly Tyr Ser Asn Glu Ile Asn Met Asp
 180 185 190

Asn Lys Pro Gly Asp Ile Ser Thr Ile Asp Gln Glu Cys Val Ser Phe
 195 200 205

Glu Thr Phe Gly Ala Asp Ile Asp Glu Ser Phe Trp Lys Glu Thr Leu
 210 215 220

Tyr Ser Gln Asp Glu His Asn Tyr Val Ser Asn Asp Leu Glu Val Ala
 225 230 235 240

Gly Leu Val Glu Ile Gln Gln Glu Phe Gln Asn Leu Gly Ser Ala Asn
 245 250 255

Asn Glu Met Ile Phe Asp Ser Glu Met Glu Leu Leu Val Arg Cys Ile
 260 265 270

Gly

<210> 41
 <211> 1262
 <212> DNA
 <213> Arabidopsis thaliana

<220>
 <221> CDS
 <222> (217)..(957)
 <223> G463

<400> 41
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 tctttctttc ttgttcttcc ttcccagggt tggttttttt tgctctctct gccttcttga 120
 ctttcaaaag actctttctt tctttggat tgattttgga ttcttaggct ctctttcttt 180

MBI-17 Sequence Listing.ST25

tagtgggttt ttgttgttgt tggttggtc tctctg atg att act gaa ctt gag Met Ile Thr Glu Leu Glu	234
1 5	
atg ggg aaa ggt gag agt gag ctt gag ctt ggt cta ggg ctg agt ctt Met Gly Lys Gly Ser Glu Leu Glu Leu Gly Leu Gly Leu Ser Leu	282
10 15 20	
ggc ggt gga acg gcg gcc aag att ggt aaa tca ggt ggt ggt ggc gcg Gly Gly Thr Ala Ala Lys Ile Gly Lys Ser Gly Gly Gly Ala	330
25 30 35	
tgg gga gag cgt gga agg ctt ttg acg gct aag gat ttt cct tct gtt Trp Gly Glu Arg Gly Arg Leu Leu Thr Ala Lys Asp Phe Pro Ser Val	378
40 45 50	
ggt tct aaa cgt gct gat tct gct tct cat gct ggt tca tct cct Gly Ser Lys Arg Ala Ala Asp Ser Ala Ser His Ala Gly Ser Ser Pro	426
55 60 65 70	
cct cgt tca agt caa gtt gtt gga tgg cct cct ata ggg tca cac agg Pro Arg Ser Ser Gln Val Val Gly Trp Pro Pro Ile Gly Ser His Arg	474
75 80 85	
atg aac agt ttg gtt aat aac caa gct aca aag tca gca aga gaa gaa Met Asn Ser Leu Val Asn Asn Gln Ala Thr Lys Ser Ala Arg Glu Glu	522
90 95 100	
gaa gaa gct ggt aag aag aaa gtg aaa gat gat gaa cct aaa gat gtg Glu Glu Ala Gly Lys Lys Val Lys Asp Asp Glu Pro Lys Asp Val	570
105 110 115	
aca aag aaa gtg aat ggg aaa gta caa gtt gga ttt att aag gtg aac Thr Lys Lys Val Asn Gly Lys Val Gln Val Gly Phe Ile Lys Val Asn	618
120 125 130	
atg gat gga gtt gct ata gga aga aaa gtg gat ttg aat gct cat tct Met Asp Gly Val Ala Ile Gly Arg Lys Val Asp Leu Asn Ala His Ser	666
135 140 145 150	
tct tac gag aat ttg gcg caa aca ttg gaa gat atg ttc ttt cgc act Ser Tyr Glu Asn Leu Ala Gln Thr Leu Glu Asp Met Phe Phe Arg Thr	714
155 160 165	
aat ccg ggt act gtc ggg tta acc agt cag ttc act aaa ccg ttg agg Asn Pro Gly Thr Val Gly Leu Thr Ser Gln Phe Thr Lys Pro Leu Arg	762
170 175 180	
ctt tta gat gga tcg tct gag ttt gta ctt act tat gaa gat aag gaa Leu Leu Asp Gly Ser Ser Glu Phe Val Leu Thr Tyr Glu Asp Lys Glu	810
185 190 195	
gga gat tgg atg ctt gtt ggt gat gtt cca tgg aga atg ttc atc aac Gly Asp Trp Met Leu Val Gly Asp Val Pro Trp Arg Met Phe Ile Asn	858
200 205 210	
tcg gtg aaa agg cta cgt gtg atg aaa acc tct gaa gct aat gga ctc Ser Val Lys Arg Leu Arg Val Met Lys Thr Ser Glu Ala Asn Gly Leu	906
215 220 225 230	
gct gca cga aat caa gaa cca aac gag aga cag cga aag cag ccg gtt Ala Ala Arg Asn Gln Glu Pro Asn Glu Arg Gln Arg Lys Gln Pro Val	954
235 240 245	
tag atcttttc gacgttacgg tgttacaggt tttatatttt ggggttttgc	1007
aagtctgaga tacttctgaa gcaaggataa gctagattga tcttatatcc agtttgtgta	1067
ttttcttggt tcttataatg gttttactg gttttcttta gtttttttt ttgctgtctt	1127
ttaattttcg gttgcgattt cactatatac tatggatgga agagaatgct ctttatatct	1187
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MBI-17 Sequence Listing.ST25

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Ser Gly Gly Gly Ala Trp Gly Glu Arg Gly Arg Leu Leu Thr Ala
35 40 45

Lys Asp Phe Pro Ser Val Gly Ser Lys Arg Ala Ala Asp Ser Ala Ser
50 55 60

His Ala Gly Ser Ser Pro Pro Arg Ser Ser Gln Val Val Gly Trp Pro
65 70 75 80

Pro Ile Gly Ser His Arg Met Asn Ser Leu Val Asn Asn Gln Ala Thr
85 90 95

Lys Ser Ala Arg Glu Glu Glu Ala Gly Lys Lys Lys Val Lys Asp
100 105 110

Asp Glu Pro Lys Asp Val Thr Lys Lys Val Asn Gly Lys Val Gln Val
115 120 125

Gly Phe Ile Lys Val Asn Met Asp Gly Val Ala Ile Gly Arg Lys Val
130 135 140

Asp Leu Asn Ala His Ser Ser Tyr Glu Asn Leu Ala Gln Thr Leu Glu
145 150 155 160

Asp Met Phe Phe Arg Thr Asn Pro Gly Thr Val Gly Leu Thr Ser Gln
165 170 175

Phe Thr Lys Pro Leu Arg Leu Leu Asp Gly Ser Ser Glu Phe Val Leu
180 185 190

Thr Tyr Glu Asp Lys Glu Gly Asp Trp Met Leu Val Gly Asp Val Pro
195 200 205

Trp Arg Met Phe Ile Asn Ser Val Lys Arg Leu Arg Val Met Lys Thr
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Gln Arg Lys Gln Pro Val
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MBI-17 Sequence Listing.ST25

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Glu Asp Ile Leu Leu Arg Gln Cys Ile Asp Lys Tyr Gly Glu Gly Lys		
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tgg cat cga gtt cct tta aga act ggt ctc aat cgg tgc cga aag agt	144	
Trp His Arg Val Pro Leu Arg Thr Gly Leu Asn Arg Cys Arg Lys Ser		
35 40 45		
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Cys Arg Leu Arg Trp Leu Asn Tyr Leu Lys Pro Ser Ile Lys Arg Gly		
50 55 60		
aaa ctc tgc tcc gat gaa gtt gat ctt gtt ctt cgc ctt cat aaa ctt	240	
Lys Leu Cys Ser Asp Glu Val Asp Leu Val Leu Arg Leu His Lys Leu		
65 70 75 80		
cta gga aat agg tgg tcc ttg atc gct ggt aga ttg cct ggt cgg act	288	
Leu Gly Asn Arg Trp Ser Leu Ile Ala Gly Arg Leu Pro Gly Arg Thr		
85 90 95		
gct aat gat gtc aag aat tac tgg aac act cat ttg agt aag aag cac	336	
Ala Asn Asp Val Lys Asn Tyr Trp Asn Thr His Leu Ser Lys Lys His		
100 105 110		
gat gaa cga tgc tgt aag acg aag atg ata aac aaa aac att act tct	384	
Asp Glu Arg Cys Cys Lys Thr Lys Met Ile Asn Lys Asn Ile Thr Ser		
115 120 125		
cat cct act tca tcg gcc caa aaa atc gat gtt tta aag cct cgg cct	432	
His Pro Thr Ser Ser Ala Gln Lys Ile Asp Val Leu Lys Pro Arg Pro		
130 135 140		
cga tcc ttc tcc gat aaa aat agt tgc aac gat gtc aat atc ttg cca	480	
Arg Ser Phe Ser Asp Lys Asn Ser Cys Asn Asp Val Asn Ile Leu Pro		
145 150 155 160		
aaa gtt gac gtt gtt cct tta cat ctt gga ctc aac aac aat tat gtt	528	
Lys Val Asp Val Val Pro Leu His Leu Gly Leu Asn Asn Tyr Val		
165 170 175		
tgt gaa agt agt att aca tgt aac aaa gat gag caa aaa gat aag ctt	576	
Cys Glu Ser Ser Ile Thr Cys Asn Lys Asp Glu Gln Lys Asp Lys Leu		
180 185 190		
att aat att aat cta ttg gat gga gat aat atg tgg tgg gaa agt tta	624	
Ile Asn Ile Asn Leu Leu Asp Gly Asp Asn Met Trp Trp Glu Ser Leu		
195 200 205		
ctg gag gca gat gtg ttg ggt cca gaa gct acg gaa aca gca aag ggt	672	
Leu Glu Ala Asp Val Leu Gly Pro Glu Ala Thr Glu Thr Ala Lys Gly		
210 215 220		
gtg acc tta ccg ctt gac ttt gag caa att tgg gct cgg ttt gat gaa	720	
Val Thr Leu Pro Leu Asp Phe Glu Gln Ile Trp Ala Arg Phe Asp Glu		
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gag act tta gaa ctg aat tag	741	
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MBI-17 Sequence Listing ST25

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 20 25 30

Trp His Arg Val Pro Leu Arg Thr Gly Leu Asn Arg Cys Arg Lys Ser
 35 40 45

Cys Arg Leu Arg Trp Leu Asn Tyr Leu Lys Pro Ser Ile Lys Arg Gly
 50 55 60

Lys Leu Cys Ser Asp Glu Val Asp Leu Val Leu Arg Leu His Lys Leu
 65 70 75 80

Leu Gly Asn Arg Trp Ser Leu Ile Ala Gly Arg Leu Pro Gly Arg Thr
 85 90 95

Ala Asn Asp Val Lys Asn Tyr Trp Asn Thr His Leu Ser Lys Lys His
 100 105 110

Asp Glu Arg Cys Cys Lys Thr Lys Met Ile Asn Lys Asn Ile Thr Ser
 115 120 125

His Pro Thr Ser Ser Ala Gln Lys Ile Asp Val Leu Lys Pro Arg Pro
 130 135 140

Arg Ser Phe Ser Asp Lys Asn Ser Cys Asn Asp Val Asn Ile Leu Pro
 145 150 155 160

Lys Val Asp Val Val Pro Leu His Leu Gly Leu Asn Asn Tyr Val
 165 170 175

Cys Glu Ser Ser Ile Thr Cys Asn Lys Asp Glu Gln Lys Asp Lys Leu
 180 185 190

Ile Asn Ile Asn Leu Leu Asp Gly Asp Asn Met Trp Trp Glu Ser Leu
 195 200 205

Leu Glu Ala Asp Val Leu Gly Pro Glu Ala Thr Glu Thr Ala Lys Gly
 210 215 220

Val Thr Leu Pro Leu Asp Phe Glu Gln Ile Trp Ala Arg Phe Asp Glu
 225 230 235 240

Glu Thr Leu Glu Leu Asn
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MBI-17 Sequence Listing.ST25

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<223> G2421

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gaa gat agt ctc ttg agg cag tgt att ggt aag tat gga gaa ggc aaa
 Glu Asp Ser Leu Leu Arg Gln Cys Ile Gly Lys Tyr Gly Glu Gly Lys
 20 25 30

96

tgg cat caa gtt cct tta aga gct ggg cta aat cgg tgc agg aaa agt
 Trp His Gln Val Pro Leu Arg Ala Gly Leu Asn Arg Cys Arg Lys Ser
 35 40 45

144

tgt aga cta aga tgg tta aac tat ttg aag cca agt atc aag aga gga
 Cys Arg Leu Arg Trp Leu Asn Tyr Leu Lys Pro Ser Ile Lys Arg Gly
 50 55 60

192

aaa ttt agt tct gat gaa gtt gat ctt ctt ctt cgt ctt cat aag ctt
 Lys Phe Ser Ser Asp Glu Val Asp Leu Leu Leu Arg Leu His Lys Leu
 65 70 75 80

240

cta gga aat agg tgg tcc ttg att gct ggt cga tta cct ggt cgg acc
 Leu Gly Asn Arg Trp Ser Leu Ile Ala Gly Arg Leu Pro Gly Arg Thr
 85 90 95

288

gct aat gat gtc aag aac tac tgg aac acc cat ctg agt aag aag cat
 Ala Asn Asp Val Lys Asn Tyr Trp Asn Thr His Leu Ser Lys Lys His
 100 105 110

336

gaa ccg tgt tgt aaa act aag ata aaa agg ata aat att ata acc cct
 Glu Pro Cys Cys Lys Thr Lys Ile Lys Arg Ile Asn Ile Ile Thr Pro
 115 120 125

384

cct aat aca ccg gcc caa aaa gtt tgt gaa aat agt atc aca tgt aac
 Pro Asn Thr Pro Ala Gln Lys Val Cys Glu Asn Ser Ile Thr Cys Asn
 130 135 140

432

aaa gat gat gag aaa gat gat ttt gtg gat aat ttt atg gtt gga gat
 Lys Asp Asp Glu Lys Asp Asp Phe Val Asp Asn Phe Met Val Gly Asp
 145 150 155 160

480

aat ata tgg ttg gag cgt ttg cta gac gag ggc caa gag gta gat gtg
 Asn Ile Trp Leu Glu Arg Leu Leu Asp Glu Gly Gln Glu Val Asp Val
 165 170 175

528

ctg gtt aca gaa gcg gcg gca aca gaa aag gag ggc act ttg gcg ttt
 Leu Val Thr Glu Ala Ala Ala Thr Glu Lys Glu Gly Thr Leu Ala Phe
 180 185 190

576

gac gtt gag caa ctt tgg aat ttg ttc gat gga gag act gtg atc ttt
 Asp Val Glu Gln Leu Trp Asn Leu Phe Asp Gly Glu Thr Val Ile Phe
 195 200 205

624

gat tag tgttataaaa cgtttgtt ctcttgtttg tgaggttct ctatthaatt
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680

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MBI-17 Sequence Listing.ST25

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35 40 45

Cys Arg Leu Arg Trp Leu Asn Tyr Leu Lys Pro Ser Ile Lys Arg Gly
50 55 60

Lys Phe Ser Ser Asp Glu Val Asp Leu Leu Leu Arg Leu His Lys Leu
65 70 75 80

Leu Gly Asn Arg Trp Ser Leu Ile Ala Gly Arg Leu Pro Gly Arg Thr
85 90 95

Ala Asn Asp Val Lys Asn Tyr Trp Asn Thr His Leu Ser Lys Lys His
 100 . . . 105 . . . 110

Glu Pro Cys Cys Lys Thr Lys Ile Lys Arg Ile Asn Ile Ile Thr Pro
115 120 125

Pro Asn Thr Pro Ala Gln Lys Val Cys Glu Asn Ser Ile Thr Cys Asn
130 135 140

Lys Asp Asp Glu Lys Asp Asp Phe Val Asp Asn Phe Met Val Gly Asp
145 150 155 160

Asn Ile Trp Leu Glu Arg Leu Leu Asp Glu Gly Gln Glu Val Asp Val
165 170 175

Leu Val Thr Glu Ala Ala Ala Thr Glu Lys Glu Gly Thr Leu Ala Phe
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195 200 205

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MBI-17 Sequence Listing ST25

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25	30	35
ctc gtg agc tat tac ttg aag agg aag gtt ctg ggt aaa cct gta cgc Leu Val Ser Tyr Tyr Leu Lys Arg Lys Val Leu Gly Lys Pro Val Arg		
40	45	55
ttc gat gcg att gga gag gta gat atc tac aag cat gag ccc tgg gat Phe Asp Ala Ile Gly Glu Val Asp Ile Tyr Lys His Glu Pro Trp Asp		
60	65	70
tta gca gtg ttt tcg aag ttg aaa act cgg gac caa gaa tgg tac ttc Leu Ala Val Phe Ser Lys Leu Lys Thr Arg Asp Gln Glu Trp Tyr Phe		
75	80	85
ttc agt gcg tta gat aag aag tac ggt aat ggt gct agg atg aat cga Phe Ser Ala Leu Asp Lys Lys Tyr Gly Asn Gly Ala Arg Met Asn Arg		
90	95	100
gca act aac aaa ggg tac tgg aaa gca act gga aaa gac aga gaa atc Ala Thr Asn Lys Gly Tyr Trp Lys Ala Thr Gly Lys Asp Arg Glu Ile		
105	110	115
cgc cgg gat att cag ttg ctc ggt atg aaa aag acg ctt gtt ttc cac Arg Arg Asp Ile Gln Leu Leu Gly Met Lys Lys Thr Leu Val Phe His		
120	125	130
agc ggg cgt gct cca gac ggc ctt cgg act aat tgg gtc atg cac gag Ser Gly Arg Ala Pro Asp Gly Leu Arg Thr Asn Trp Val Met His Glu		
140	145	150
tat cgc ctt gtg gaa tat gaa act gaa act aac gga agc ctg ctg cag Tyr Arg Leu Val Glu Tyr Glu Thr Glu Thr Asn Gly Ser Leu Leu Gln		
155	160	165
gat gca tat gtg ttg tgc aga qtg ttt cac aag aat aac att ggg cca Asp Ala Tyr Val Leu Cys Arg Val Phe His Lys Asn Asn Ile Gly Pro		
170	175	180
cca agt ggg aac aga tat gcg cca ttc atg gaa gaa gaa tgg gct gat Pro Ser Gly Asn Arg Tyr Ala Pro Phe Met Glu Glu Trp Ala Asp		
185	190	195
ggg gga gga gct ctg att cca gga ata gac gtt agg gtc agg gta gag Gly Gly Gly Ala Leu Ile Pro Gly Ile Asp Val Arg Val Arg Val Glu		
200	205	210
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220	225	230
tca gca agc aag gat ctc att aac atc aac gag cta ccg aga gat gct Ser Ala Ser Lys Asp Leu Ile Asn Ile Asn Glu Leu Pro Arg Asp Ala		
235	240	245
act cca atg gac atc gaa cct aac caa cag aat cat cat gag agt gcc Thr Pro Met Asp Ile Glu Pro Asn Gln Gln Asn His His Glu Ser Ala		
250	255	260
ttc aag cca cag gag agt aac aac cat agt ggt tat gaa gaa gat gag Phe Lys Pro Gln Glu Ser Asn Asn His Ser Gly Tyr Glu Asp Glu		
265	270	275
gac aca ctc aaa cgc gag cac gca gaa gaa gat gag cgt cct cct tct Asp Thr Leu Lys Arg Glu His Ala Glu Glu Asp Glu Arg Pro Pro Ser		
280	285	295
cta tgc att ctc aac aaa gaa gct cca cta cct ctc ctg caa tac aaa Leu Cys Ile Leu Asn Lys Glu Ala Pro Leu Pro Leu Leu Gln Tyr Lys		
300	305	310

MBI-17 Sequence Listing ST25

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atc tca tca tct gct gct gct acc aac act gcc atc tct gca ttg Ile Ser Ser Ala Ala Ala Ala Thr Asn Thr Ala Ile Ser Ala Leu 345 350 355	1109
ctt gag ttc tca ctt atg ggt atc tcc gac aag aaa gaa aac cag cag Leu Glu Phe Ser Leu Met Gly Ile Ser Asp Lys Lys Glu Asn Gln Gln 360 365 370 375	1157
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gtt aat gat ctc cag aag gag gtt cac cag atg tct gtt gaa aga gaa Val Asn Asp Leu Gln Lys Glu Val His Gln Met Ser Val Glu Arg Glu 395 400 405	1253
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cag tca aga atc gat gcg ctg cgt cag gag aac gag gaa ctt aag aag Gln Ser Arg Ile Asp Ala Leu Arg Gln Glu Asn Glu Glu Leu Lys Lys 425 430 435	1349
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Phe His Pro Thr Asp Glu Glu Leu Val Ser Tyr Tyr Leu Lys Arg Lys 35 40 45	
Val Leu Gly Lys Pro Val Arg Phe Asp Ala Ile Gly Glu Val Asp Ile 50 55 60	
Tyr Lys His Glu Pro Trp Asp Leu Ala Val Phe Ser Lys Leu Lys Thr 65 70 75 80	

Arg Asp Gln Glu Trp Tyr Phe Phe Ser Ala Leu Asp Lys Lys Tyr Gly

MBI-17 Sequence Listing ST25

85 90 95

Asn Gly Ala Arg Met Asn Arg Ala Thr Asn Lys Gly Tyr Trp Lys Ala
 100 105 110

Thr Gly Lys Asp Arg Glu Ile Arg Arg Asp Ile Gln Leu Leu Gly Met
 115 120 125

Lys Lys Thr Leu Val Phe His Ser Gly Arg Ala Pro Asp Gly Leu Arg
 130 135 140

Thr Asn Trp Val Met His Glu Tyr Arg Leu Val Glu Tyr Glu Thr Glu
 145 150 155 160

Thr Asn Gly Ser Leu Leu Gln Asp Ala Tyr Val Leu Cys Arg Val Phe
 165 170 175

His Lys Asn Asn Ile Gly Pro Pro Ser Gly Asn Arg Tyr Ala Pro Phe
 180 185 190

Met Glu Glu Glu Trp Ala Asp Gly Gly Ala Leu Ile Pro Gly Ile
 195 200 205

Asp Val Arg Val Arg Val Glu Ala Leu Pro Gln Ala Asn Gly Asn Asn
 210 215 220

Gln Met Asp Gln Glu Met His Ser Ala Ser Lys Asp Leu Ile Asn Ile
 225 230 235 240

Asn Glu Leu Pro Arg Asp Ala Thr Pro Met Asp Ile Glu Pro Asn Gln
 245 250 255

Gln Asn His His Glu Ser Ala Phe Lys Pro Gln Glu Ser Asn Asn His
 260 265 270

Ser Gly Tyr Glu Glu Asp Glu Asp Thr Leu Lys Arg Glu His Ala Glu
 275 280 285

Glu Asp Glu Arg Pro Pro Ser Leu Cys Ile Leu Asn Lys Glu Ala Pro
 290 295 300

Leu Pro Leu Leu Gln Tyr Lys Arg Arg Arg Gln Asn Glu Ser Asn Asn
 305 310 315 320

Asn Ser Ser Arg Asn Thr Gln Asp His Cys Ser Ser Thr Ile Thr Thr
 325 330 335

Val Asp Asn Thr Thr Leu Ile Ser Ser Ser Ala Ala Ala Thr
 340 345 350

Asn Thr Ala Ile Ser Ala Leu Leu Glu Phe Ser Leu Met Gly Ile Ser
 355 360 365

Asp Lys Lys Glu Asn Gln Gln Lys Glu Glu Thr Ser Pro Pro Ser Pro
 370 375 380

MBI-17 Sequence Listing.ST25

Ile Ala Ser Pro Glu Glu Lys Val Asn Asp Leu Gln Lys Glu Val His
 385 390 395 400

Gln Met Ser Val Glu Arg Glu Thr Phe Lys Leu Glu Met Met Ser Ala
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Glu Ala Met Ile Ser Ile Leu Gln Ser Arg Ile Asp Ala Leu Arg Gln
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Glu Asn Glu Glu Leu Lys Lys Asn Ala Ser Gly Gln Ala Ser
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Thr Val Asp Ile Met Arg Leu Pro Lys Met Glu Asp Gln Thr Ala Ile
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caa gaa gct gca tca caa ggc tta aaa agc atg gaa cac ttg att cgt 154
Gln Glu Ala Ala Ser Gln Gly Leu Lys Ser Met Glu His Leu Ile Arg
20 25 30

gtc ctc tct aac cgt ccc gaa gaa cgt aac gtt gat tgc tct gag atc 202
Val Leu Ser Asn Arg Pro Glu Glu Arg Asn Val Asp Cys Ser Glu Ile
35 40 45

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Thr Asp Phe Thr Val Ser Lys Phe Lys Val Ile Ser Leu Leu Asn
50 55 60 65

cgt tcc ggt cac gcc cgg ttt aga cgt ggt ccg gtt cat tcc cct cct 298
Arg Ser Gly His Ala Arg Phe Arg Arg Gly Pro Val His Ser Pro Pro
70 75 80

tcc tcc tcc gtt cct cca ccg gtg aaa gtg aca act ccg gct ccc act 346
Ser Ser Ser Val Pro Pro Pro Val Lys Val Thr Thr Pro Ala Pro Thr
85 90 95

cag atc tct gct cca gca ccg gtt agc ttc gtt cag gca aat caa caa 394
Gln Ile Ser Ala Pro Ala Pro Val Ser Phe Val Gln Ala Asn Gln Gln
100 105 110

agc gtg acg tta gat ttc act aga ccg agc gtt ttt ggc gct aaa acc 442
Ser Val Thr Leu Asp Phe Thr Arg Pro Ser Val Phe Gly Ala Lys Thr
115 120 125

aag agc tcg gag gtt gag ttt gct aaa gag agc ttt agc gta tct 490
Lys Ser Ser Glu Val Val Glu Phe Ala Lys Glu Ser Phe Ser Val Ser
130 135 140 145

tct aac tct tct ttc atg tct tct gcg atc acc ggt gat gga agt gtc 538
Ser Asn Ser Ser Phe Met Ser Ser Ala Ile Thr Gly Asp Gly Ser Val
150 155 160

tct aaa ggc tct tcg atc ttt ctt gct ccg gct cca gcg gtg cca gtg 586
Ser Lys Gly Ser Ser Ile Phe Leu Ala Pro Ala Pro Ala Val Pro Val
165 170 175

MBI-17 Sequence Listing.ST25

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Cys Phe His Asp His Ser Glu Gly Phe Ser Gly Lys Ile Ser Gly	
195 200 205	
tcc ggc aac ggc aag tgc cat tgc aag aaa agc cga aaa aat cgg atg	730
Ser Gly Asn Gly Lys Cys His Cys Lys Ser Arg Lys Asn Arg Met	
210 215 220 225	
aag aga acc gtg aga gta ccg gcg gta agt gca aag atc gcc gat ata	778
Lys Arg Thr Val Arg Val Pro Ala Val Ser Ala Lys Ile Ala Asp Ile	
230 235 240	
cca cca gac gaa tat tca tgg aga aag tat gga caa aaa ccg atc aaa	826
Pro Pro Asp Glu Tyr Ser Trp Arg Lys Tyr Gly Gln Lys Pro Ile Lys	
245 250 255	
ggc tca cca cat cca ccg ggt tat tac aag tgt agt aca ttt aga gga	874
Gly Ser Pro His Pro Arg Gly Tyr Tyr Lys Cys Ser Thr Phe Arg Gly	
260 265 270	
tgt cca gcg agg aaa cac gtg gaa aga gct ttg gat gat tca acg atg	922
Cys Pro Ala Arg Lys His Val Glu Arg Ala Leu Asp Asp Ser Thr Met	
275 280 285	
ttg att gtg acg tac gaa gga gag cac cgt cat cac cag tcc acg atg	970
Leu Ile Val Thr Tyr Glu Gly His Arg His His Gln Ser Thr Met	
290 295 300 305	
cag gag cat gta act cct agc gtg agt ggt ttg gtg ttt ggt tcg gct	1018
Gln Glu His Val Thr Pro Ser Val Ser Gly Leu Val Phe Gly Ser Ala	
310 315 320	
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ggttttgtaa tttttttct ataacaaaat tagttttaga tttttttta gtatgttttt	1131
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Arg Val Leu Ser Asn Arg Pro Glu Glu Arg Asn Val Asp Cys Ser Glu	
35 40 45	
Ile Thr Asp Phe Thr Val Ser Lys Phe Lys Lys Val Ile Ser Leu Leu	
50 55 60	
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Pro Ser Ser Ser Val Pro Pro Val Lys Val Thr Thr Pro Ala Pro	
85 90 95	

MBI-17 Sequence Listing.ST25

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 100 105 110

Gln Ser Val Thr Leu Asp Phe Thr Arg Pro Ser Val Phe Gly Ala Lys
 115 120 125

Thr Lys Ser Ser Glu Val Val Glu Phe Ala Lys Glu Ser Phe Ser Val
 130 135 140

Ser Ser Asn Ser Ser Phe Met Ser Ser Ala Ile Thr Gly Asp Gly Ser
 145 150 155 160

Val Ser Lys Gly Ser Ser Ile Phe Leu Ala Pro Ala Pro Ala Val Pro
 165 170 175

Val Thr Ser Ser Gly Lys Pro Pro Leu Ser Gly Leu Pro Tyr Arg Lys
 180 185 190

Arg Cys Phe Glu His Asp His Ser Glu Gly Phe Ser Gly Lys Ile Ser
 195 200 205

Gly Ser Gly Asn Gly Lys Cys His Cys Lys Ser Arg Lys Asn Arg
 210 215 220

Met Lys Arg Thr Val Arg Val Pro Ala Val Ser Ala Lys Ile Ala Asp
 225 230 235 240

Ile Pro Pro Asp Glu Tyr Ser Trp Arg Lys Tyr Gly Gln Lys Pro Ile
 245 250 255

Lys Gly Ser Pro His Pro Arg Gly Tyr Tyr Lys Cys Ser Thr Phe Arg
 260 265 270

Gly Cys Pro Ala Arg Lys His Val Glu Arg Ala Leu Asp Asp Ser Thr
 275 280 285

Met Leu Ile Val Thr Tyr Glu Gly Glu His Arg His His Gln Ser Thr
 290 295 300

Met Gln Glu His Val Thr Pro Ser Val Ser Gly Leu Val Phe Gly Ser
 305 310 315 320

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atg atg ttt	aat gag atg	gga atg tgt	gga aac atg	gat ttc ttc	tct	226
Met Met Phe	Asn Glu Met	Gly Met Cys	Gly Asn Met	Asp Phe Phe	Ser	
1	5	10	15			
tct gga tca ctt	ggt gaa gtt gat	ttc tgt cct	gtt cca caa	gct gag		274
Ser Gly Ser	Leu Gly Glu Val	Asp Phe Cys Pro	Val Pro Gln	Ala Glu		
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cct gat tcc att	gtt gaa gat gac	tat act gat gat	gag att gat	gtt		322
Pro Asp Ser	Ile Val Glu Asp	Asp Tyr Thr Asp	Asp Glu Ile	Asp Val		
35	40	45				
gat gaa ttg gag	agg agg atg tgg	aga gac aaa	atg cgg ctt	aaa cgt		370
Asp Glu Leu	Glu Arg Arg Met Trp	Arg Asp Lys	Met Arg Leu	Lys Arg		
50	55	60				
ctc aag gag cag	gat aag ggt	aaa gaa ggt	gtt gat	gct gct	aaa cag	418
Leu Lys Glu	Gln Asp Lys	Gly Lys Glu	Gly Val	Asp Ala	Ala Lys Gln	
65	70	75	80			
agg cag tct caa	gag caa gct	agg agg aag	aaa atg tct	aga gct	caa	466
Arg Gln Ser	Gln Glu Gln Ala	Arg Arg Lys	Lys Met Ser	Arg Ala	Gln	
85	90	95				
gat ggg atc ttg	aag tat atg ttg	aag atg atg	gaa gtt	tgt aaa	gct	514
Asp Gly Ile	Leu Lys Tyr Met	Leu Lys Met	Met Glu	Val Cys	Lys Ala	
100	105	110				
caa ggc ttt gtt	tat ggg att att	ccg gag aat	ggg aag cct	gtg act		562
Gln Gly Phe	Val Tyr Ile	Ile Pro Glu Asn	Gly Lys	Pro Val Thr		
115	120	125				
ggt gct tct	aat tta agg gag	tgg tgg aaa	aat gat aag	gtt agg ttt		610
Gly Ala Ser	Asp Asn Leu	Arg Glu Trp	Trp Lys	Asp Lys Val	Arg Phe	
130	135	140				
gat cgt aat	ggt cct gcg	gct att acc aag	tat caa	gcg gag aat	aat	658
Asp Arg Asn	Gly Pro Ala	Ala Ile Thr	Lys Tyr	Gln Ala Glu	Asn Asn	
145	150	155	160			
atc ccg ggg att	cat gaa ggt	aat aac ccg att	gga ccg	act cct cat		706
Ile Pro Gly	Ile His Glu	Gly Asn Asn	Pro Ile	Gly Pro Thr	Pro His	
165	170	175				
acc ttg caa gag	ctt caa gac acg	act ctt gga tcg	ctt ttg tct	gcg		754
Thr Leu Gln	Glu Leu Gln Asp	Thr Thr Leu	Gly Ser	Leu Leu Ser	Ala	
180	185	190				
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Leu Met Gln	His Cys Asp	Pro Pro Gln	Arg Arg Phe	Pro Leu Glu	Lys	
195	200	205				
gga gtt cct cct	ccg tgg cct	aat ggg aaa	gag gat	tgg tgg cct		850
Gly Val Pro	Pro Pro Trp	Pro Asn Gly	Lys Glu Asp	Trp Trp Pro		
210	215	220				
caa ctt ggt	ttg cct aaa	aat gat caa	ggc cct tac	aag aag cct		898
Gln Leu	Gly Leu Pro	Lys Asp Gln	Gly Pro Ala	Pro Tyr Lys	Pro	
225	230	235	240			
cat gat ttg	aag aag gcg	tgg aaa gtc	ggc gtt ttg	act gcg	gtt atc	946
His Asp Leu	Lys Lys Ala	Trp Lys Val	Gly Val	Leu Thr	Ala Val Ile	
245	250	255				
aag cat atg	ttt cct gat	att gct aag atc	cgt aag ctc	gtg agg	caa	994
Lys His Met	Phe Pro Asp Ile	Ala Lys Ile	Arg Lys Leu	Val Arg	Gln	
260	265	270				
tct aaa tgg	cag gat aag atg	act gct aaa	gag agt	gct acc	tgg	1042
Ser Lys Cys	Leu Gln Asp	Lys Met Thr	Ala Lys Glu	Ser Ala	Thr Trp	

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275 280 285

ctt gct att att aac caa gag gag tcc ttg gct aga gag ctt tat ccc Leu Ala Ile Ile Asn Gln Glu Glu Ser Leu Ala Arg Glu Leu Tyr Pro 290 295 300	1090
gag tca tgt cca cct ctt tct ctg tct ggt gga agt tgc tcg ctt ctg Glu Ser Cys Pro Pro Leu Ser Leu Ser Gly Gly Ser Cys Ser Leu Leu 305 310 315 320	1138
atg aat gat tgc agt caa tac gat gtt gaa ggt ttc gag aag gag tct Met Asn Asp Cys Ser Gln Tyr Asp Val Glu Gly Phe Glu Lys Glu Ser 325 330 335	1186
cac tat gaa gtg gaa gag ctc aag cca gaa aaa gtt atg aat tct tca His Tyr Glu Val Glu Leu Lys Pro Glu Lys Val Met Asn Ser Ser 340 345 350	1234
aac ttt ggg atg gtt gct aaa atg cat gac ttt cct gtc aaa gaa gaa Asn Phe Gly Met Val Ala Lys Met His Asp Phe Pro Val Lys Glu Glu 355 360 365	1282
gtc cca gca gga aac tcg gaa ttc atg aga aag aga aag cca aac aga Val Pro Ala Gly Asn Ser Glu Phe Met Arg Lys Arg Lys Pro Asn Arg 370 375 380	1330
gat ctg aac act att atg gac aga acc gtt ttc acc tgc gag aat ctt Asp Leu Asn Thr Ile Met Asp Arg Thr Val Phe Thr Cys Glu Asn Leu 385 390 395 400	1378
ggg tgt gcg cac agc gaa atc agc cgg gga ttt ctg gat agg aat tcg Gly Cys Ala His Ser Glu Ile Ser Arg Gly Phe Leu Asp Arg Asn Ser 405 410 415	1426
aga gac aac cat caa ctg gca tgt cca cat cga gac agt cgc tta ccg Arg Asp Asn His Gln Leu Ala Cys Pro His Arg Asp Ser Arg Leu Pro 420 425 430	1474
tat gga gca gca cca agg ttt cat gtc aat gaa gtt aag cct gta Tyr Gly Ala Ala Pro Ser Arg Phe His Val Asn Glu Val Lys Pro Val 435 440 445	1522
gtt gga ttt cct cag cca agg cca gtg aac tca gta gcc caa cca att Val Gly Phe Pro Gln Pro Arg Pro Val Asn Ser Val Ala Gln Pro Ile 450 455 460	1570
gac tta acg ggt ata gtt cct gaa gat gga cag aag atg atc tca gag Asp Leu Thr Gly Ile Val Pro Glu Asp Gly Gln Lys Met Ile Ser Glu 465 470 475 480	1618
ctc atg tcc atg tac gac aga aat gtc cag agc aac caa acc tct atg Leu Met Ser Met Tyr Asp Arg Asn Val Gln Ser Asn Gln Thr Ser Met 485 490 495	1666
gtc atg gaa aat caa agc gtg tca ctg ctt caa ccc aca gtc cat aac Val Met Glu Asn Gln Ser Val Ser Leu Leu Gln Pro Thr Val His Asn 500 505 510	1714
cat caa gaa cat ctc cag ttc cca gga aac atg gtg gaa gga agt ttc His Gln Glu His Leu Gln Phe Pro Gly Asn Met Val Glu Gly Ser Phe 515 520 525	1762
ttt gaa gac ttg aac atc cca aac aga gca aac aac aac aac agc agc Phe Glu Asp Leu Asn Ile Pro Asn Arg Ala Asn Asn Asn Ser Ser 530 535 540	1810
aac aat caa acg ttt ttt caa ggg aac aac aac aac aac aat gtg ttt Asn Asn Gln Thr Phe Phe Gln Gly Asn Asn Asn Asn Asn Val Phe 545 550 555 560	1858
aag ttc gac act gca gat cac aac aac ttt gaa gct gca cat aac aac Lys Phe Asp Thr Ala Asp His Asn Asn Phe Glu Ala Ala His Asn Asn 565 570 575	1906
aac aat aac agt agc ggc aac agg ttc cag ctt gtg ttt gat tcc aca	1954

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Asn Asn Asn Ser Ser Gly Asn Arg Phe Gln Leu Val Phe Asp Ser Thr																																																																																																																																															
580	585	590				ccg ttc gac atg gcg tca ttc gat tac .aga gat gat atg tcg atg cca	2002	Pro Phe Asp Met Ala Ser Phe Asp Tyr Arg Asp Asp Met Ser Met Pro		595	600	605				gga gta gta gga acg atg gat gga atg cag cag aag cag caa gat gta	2050	Gly Val Val Gly Thr Met Asp Gly Met Gln Gln Lys Gln Gln Asp Val		610	615	620				tcc ata tgg ttc taa agtcttgta gtagatttca tcttcctta ttttatctt	2105	Ser Ile Trp Phe		625				ttgtgttctt acattcactc aaccatgtaa tatttttcc tgggtctctc tgtctctatc	2165	gcttgttatg atgtgtctgt aagagtctct aaaaactctc tgttactgtg tgtctttgtc	2225	tcggcttggt gaatctctct gtcatcatca gcttttagtt acacacccga cttggggatg	2285	aacgaacact aaatgtaaat ttca	2310			<210> 52		<211> 628		<212> PRT		<213> Arabidopsis thaliana				<400> 52				Met Met Phe Asn Glu Met Gly Met Cys Gly Asn Met Asp Phe Phe Ser		1	5	10	15			Ser Gly Ser Leu Gly Glu Val Asp Phe Cys Pro Val Pro Gln Ala Glu		20	25	30				Pro Asp Ser Ile Val Glu Asp Asp Tyr Thr Asp Asp Glu Ile Asp Val		35	40	45				Asp Glu Leu Glu Arg Arg Met Trp Arg Asp Lys Met Arg Leu Lys Arg		50	55	60				Leu Lys Glu Gln Asp Lys Gly Lys Glu Gly Val Asp Ala Ala Lys Gln		65	70	75	80			Arg Gln Ser Gln Glu Gln Ala Arg Arg Lys Lys Met Ser Arg Ala Gln		85	90	95				Asp Gly Ile Leu Lys Tyr Met Leu Lys Met Met Glu Val Cys Lys Ala		100	105	110				Gln Gly Phe Val Tyr Gly Ile Ile Pro Glu Asn Gly Lys Pro Val Thr		115	120	125				Gly Ala Ser Asp Asn Leu Arg Glu Trp Trp Lys Asp Lys Val Arg Phe		130	135	140				Asp Arg Asn Gly Pro Ala Ala Ile Thr Lys Tyr Gln Ala Glu Asn Asn		145	150	155	160			Ile Pro Gly Ile His Glu Gly Asn Asn Pro Ile Gly Pro Thr Pro His		165	170	175	
590																																																																																																																																															
ccg ttc gac atg gcg tca ttc gat tac .aga gat gat atg tcg atg cca	2002																																																																																																																																														
Pro Phe Asp Met Ala Ser Phe Asp Tyr Arg Asp Asp Met Ser Met Pro																																																																																																																																															
595	600	605				gga gta gta gga acg atg gat gga atg cag cag aag cag caa gat gta	2050	Gly Val Val Gly Thr Met Asp Gly Met Gln Gln Lys Gln Gln Asp Val		610	615	620				tcc ata tgg ttc taa agtcttgta gtagatttca tcttcctta ttttatctt	2105	Ser Ile Trp Phe		625				ttgtgttctt acattcactc aaccatgtaa tatttttcc tgggtctctc tgtctctatc	2165	gcttgttatg atgtgtctgt aagagtctct aaaaactctc tgttactgtg tgtctttgtc	2225	tcggcttggt gaatctctct gtcatcatca gcttttagtt acacacccga cttggggatg	2285	aacgaacact aaatgtaaat ttca	2310			<210> 52		<211> 628		<212> PRT		<213> Arabidopsis thaliana				<400> 52				Met Met Phe Asn Glu Met Gly Met Cys Gly Asn Met Asp Phe Phe Ser		1	5	10	15			Ser Gly Ser Leu Gly Glu Val Asp Phe Cys Pro Val Pro Gln Ala Glu		20	25	30				Pro Asp Ser Ile Val Glu Asp Asp Tyr Thr Asp Asp Glu Ile Asp Val		35	40	45				Asp Glu Leu Glu Arg Arg Met Trp Arg Asp Lys Met Arg Leu Lys Arg		50	55	60				Leu Lys Glu Gln Asp Lys Gly Lys Glu Gly Val Asp Ala Ala Lys Gln		65	70	75	80			Arg Gln Ser Gln Glu Gln Ala Arg Arg Lys Lys Met Ser Arg Ala Gln		85	90	95				Asp Gly Ile Leu Lys Tyr Met Leu Lys Met Met Glu Val Cys Lys Ala		100	105	110				Gln Gly Phe Val Tyr Gly Ile Ile Pro Glu Asn Gly Lys Pro Val Thr		115	120	125				Gly Ala Ser Asp Asn Leu Arg Glu Trp Trp Lys Asp Lys Val Arg Phe		130	135	140				Asp Arg Asn Gly Pro Ala Ala Ile Thr Lys Tyr Gln Ala Glu Asn Asn		145	150	155	160			Ile Pro Gly Ile His Glu Gly Asn Asn Pro Ile Gly Pro Thr Pro His		165	170	175											
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Gly Val Val Gly Thr Met Asp Gly Met Gln Gln Lys Gln Gln Asp Val																																																																																																																																															
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Ser Ile Trp Phe																																																																																																																																															
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1	5	10	15			Ser Gly Ser Leu Gly Glu Val Asp Phe Cys Pro Val Pro Gln Ala Glu		20	25	30				Pro Asp Ser Ile Val Glu Asp Asp Tyr Thr Asp Asp Glu Ile Asp Val		35	40	45				Asp Glu Leu Glu Arg Arg Met Trp Arg Asp Lys Met Arg Leu Lys Arg		50	55	60				Leu Lys Glu Gln Asp Lys Gly Lys Glu Gly Val Asp Ala Ala Lys Gln		65	70	75	80			Arg Gln Ser Gln Glu Gln Ala Arg Arg Lys Lys Met Ser Arg Ala Gln		85	90	95				Asp Gly Ile Leu Lys Tyr Met Leu Lys Met Met Glu Val Cys Lys Ala		100	105	110				Gln Gly Phe Val Tyr Gly Ile Ile Pro Glu Asn Gly Lys Pro Val Thr		115	120	125				Gly Ala Ser Asp Asn Leu Arg Glu Trp Trp Lys Asp Lys Val Arg Phe		130	135	140				Asp Arg Asn Gly Pro Ala Ala Ile Thr Lys Tyr Gln Ala Glu Asn Asn		145	150	155	160			Ile Pro Gly Ile His Glu Gly Asn Asn Pro Ile Gly Pro Thr Pro His		165	170	175																																																													
10	15																																																																																																																																														
Ser Gly Ser Leu Gly Glu Val Asp Phe Cys Pro Val Pro Gln Ala Glu																																																																																																																																															
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Pro Asp Ser Ile Val Glu Asp Asp Tyr Thr Asp Asp Glu Ile Asp Val																																																																																																																																															
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Arg Gln Ser Gln Glu Gln Ala Arg Arg Lys Lys Met Ser Arg Ala Gln																																																																																																																																															
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Gln Gly Phe Val Tyr Gly Ile Ile Pro Glu Asn Gly Lys Pro Val Thr																																																																																																																																															
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Asp Arg Asn Gly Pro Ala Ala Ile Thr Lys Tyr Gln Ala Glu Asn Asn																																																																																																																																															
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Ile Pro Gly Ile His Glu Gly Asn Asn Pro Ile Gly Pro Thr Pro His																																																																																																																																															
165	170	175																																																																																																																																													
175																																																																																																																																															

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Thr	Leu	Gln	Glu	Leu	Gln	Asp	Thr	Thr	Leu	Gly	Ser	Leu	Leu	Ser	Ala
180															
185															
190															
Leu	Met	Gln	His	Cys	Asp	Pro	Pro	Gln	Arg	Arg	Phe	Pro	Leu	Glu	Lys
195															
200															
205															
Gly	Val	Pro	Pro	Pro	Trp	Trp	Pro	Asn	Gly	Lys	Glu	Asp	Trp	Trp	Pro
210															
215															
220															
Gln	Leu	Gly	Leu	Pro	Lys	Asp	Gln	Gly	Pro	Ala	Pro	Tyr	Lys	Pro	
225															
230															
235															
His	Asp	Leu	Lys	Lys	Ala	Trp	Lys	Val	Gly	Val	Leu	Thr	Ala	Val	Ile
245															
250															
255															
Lys	His	Met	Phe	Pro	Asp	Ile	Ala	Lys	Ile	Arg	Lys	Leu	Val	Arg	Gln
260															
265															
270															
Ser	Lys	Cys	Leu	Gln	Asp	Lys	Met	Thr	Ala	Lys	Glu	Ser	Ala	Thr	Trp
275															
280															
285															
Leu	Ala	Ile	Ile	Asn	Gln	Glu	Glu	Ser	Leu	Ala	Arg	Glu	Leu	Tyr	Pro
290															
295															
300															
Glu	Ser	Cys	Pro	Pro	Leu	Ser	Leu	Ser	Gly	Gly	Ser	Cys	Ser	Leu	Leu
305															
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Met	Asn	Asp	Cys	Ser	Gln	Tyr	Asp	Val	Glu	Gly	Phe	Glu	Lys	Glu	Ser
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His	Tyr	Glu	Val	Glu	Glu	Leu	Lys	Pro	Glu	Lys	Val	Met	Asn	Ser	Ser
340															
345															
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Asn	Phe	Gly	Met	Val	Ala	Lys	Met	His	Asp	Phe	Pro	Val	Lys	Glu	Glu
355															
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365															
Val	Pro	Ala	Gly	Asn	Ser	Glu	Phe	Met	Arg	Lys	Arg	Lys	Pro	Asn	Arg
370															
375															
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Asp	Leu	Asn	Thr	Ile	Met	Asp	Arg	Thr	Val	Phe	Thr	Cys	Glu	Asn	Leu
385															
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395															
Gly	Cys	Ala	His	Ser	Glu	Ile	Ser	Arg	Gly	Phe	Leu	Asp	Arg	Asn	Ser
405															
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Arg	Asp	Asn	His	Gln	Leu	Ala	Cys	Pro	His	Arg	Asp	Ser	Arg	Leu	Pro
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430															
Tyr	Gly	Ala	Ala	Pro	Ser	Arg	Phe	His	Val	Asn	Glu	Val	Lys	Pro	Val
435															
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Val	Gly	Phe	Pro	Gln	Pro	Arg	Pro	Val	Asn	Ser	Val	Ala	Gln	Pro	Ile
450															
455															
460															
Asp	Leu	Thr	Gly	Ile	Val	Pro	Glu	Asp	Gly	Gln	Lys	Met	Ile	Ser	Glu
465															
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475															

MBI-17 Sequence Listing.ST25

Leu Met Ser Met Tyr Asp Arg Asn Val Gln Ser Asn Gln Thr Ser Met
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Val Met Glu Asn Gln Ser Val Ser Leu Leu Gln Pro Thr Val His Asn
500 505 510

His Gln Glu His Leu Gln Phe Pro Gly Asn Met Val Glu Gly Ser Phe
515 520 525

Phe Glu Asp Leu Asn Ile Pro Asn Arg Ala Asn Asn Asn Ser Ser
530 535 540

Asn Asn Gln Thr Phe Phe Gln Gly Asn Asn Asn Asn Asn Val Phe
545 550 555 560

Lys Phe Asp Thr Ala Asp His Asn Asn Phe Glu Ala Ala His Asn Asn
565 570 575

Asn Asn Asn Ser Ser Gly Asn Arg Phe Gln Leu Val Phe Asp Ser Thr
580 585 590

Pro Phe Asp Met Ala Ser Phe Asp Tyr Arg Asp Asp Met Ser Met Pro
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Ser Ile Trp Phe
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tgg aca tct gaa gaa gac cag aag ctt gtt gac tat atc cag aaa cat 96
Trp Thr Ser Glu Glu Asp Gln Lys Leu Val Asp Tyr Ile Gln Lys His
20 25 30

ggt tat ggt aac tgg aga acc ctc ccc aaa aat gcc ggt acg tgt ttg 144
Gly Tyr Gly Asn Trp Arg Thr Leu Pro Lys Asn Ala Gly Thr Cys Leu
35 40 45

caa aga tgt ggc aaa agt tgt agg tta agg tgg act aat tat ctc cga 192
Gln Arg Cys Gly Lys Ser Cys Arg Leu Arg Trp Thr Asn Tyr Leu Arg
50 55 60

cca gat ata aaa cga gga aga ttc tct ttt gag gaa gaa gaa gcc att 240
Pro Asp Ile Lys Arg Gly Arg Phe Ser Phe Glu Glu Glu Ala Ile
65 70 75 80

att cag ctt cat agc ttc tta gga aac aag tgg tct gcg att gcg gcg 288
Ile Gln Leu His Ser Phe Leu Gly Asn Lys Trp Ser Ala Ile Ala Ala

MBI-17 Sequence Listing ST25

85 90 95

cgt ttg cca gga aga aca gat aat gag atc aag aac ttt tgg aac act Arg Leu Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Phe Trp Asn Thr 100 105 110	336
cat ata aga aag aag cta ctt aga atg ggg att gat cca gtg act cac His Ile Arg Lys Lys Leu Leu Arg Met Gly Ile Asp Pro Val Thr His 115 120 125	384
agt cca cga ctc gat ctc ctc gat atc tca tcc atc tta gct tca tct Ser Pro Arg Leu Asp Leu Asp Ile Ser Ser Ile Leu Ala Ser Ser 130 135 140	432
cta tac aat tca tct tca cat cac atg aac atg tca aga ctc atg atg Leu Tyr Asn Ser Ser His His Met Asn Met Ser Arg Leu Met Met 145 150 155 160	480
gat act aat cgt cgt cat cag caa caa cat cca ttg gtt aac ccc gag Asp Thr Asn Arg Arg His Gln Gln His Pro Leu Val Asn Pro Glu 165 170 175	528
ata ctc aag ctt gcg acc tct ata ttc tct caa aac caa aac caa aac Ile Leu Lys Leu Ala Thr Ser Ile Phe Ser Gln Asn Gln Asn Gln Asn 180 185 190	576
cac aac caa aat caa aac caa aac caa aac ctc gtg gtg gat cat gag His Asn Gln Asn Gln Asn Gln Asn Leu Val Val Asp His Glu 195 200 205	624
aag caa aca gtt tat cat cat cat gat gtt aac caa acc gga gta aac Lys Gln Thr Val Tyr His His Asp Val Asn Gln Thr Gly Val Asn 210 215 220	672
caa tac caa acc gac caa tat ttc gag aac gcg att act caa gaa ctc Gln Tyr Gln Thr Asp Gln Tyr Phe Glu Asn Ala Ile Thr Gln Glu Leu 225 230 235 240	720
caa tct tcc atg cca cca ttc ccc aat gaa gct cat cag ttt aac gac Gln Ser Ser Met Pro Pro Phe Pro Asn Glu Ala His Gln Phe Asn Asp 245 250 255	768
atg gat cat cac ttc aat ggt ttt gga gaa caa aat ctt gtt tca act Met Asp His His Phe Asn Gly Phe Gly Glu Gln Asn Leu Val Ser Thr 260 265 270	816
tct act acg tca gtc caa gat tgc tat aat ccg tca ttc aac gat tat Ser Thr Thr Ser Val Gln Asp Cys Tyr Asn Pro Ser Phe Asn Asp Tyr 275 280 285	864
tca agt tca aat ttt gtc tta gat cat tct tat tcg gat cag agc ttc Ser Ser Ser Asn Phe Val Leu Asp His Ser Tyr Ser Asp Gln Ser Phe 290 295 300	912
aac ttc gca aat tcg gtc tta aac acg cca tcc tcg agc ccg agc ccg Asn Phe Ala Asn Ser Val Leu Asn Thr Pro Ser Ser Ser Pro Ser Pro 305 310 315 320	960
act acg tta aac tcg agt tac atc aat agt agc agt tgc agc act gag Thr Thr Leu Asn Ser Ser Tyr Ile Asn Ser Ser Ser Cys Ser Thr Glu 325 330 335	1008
gat gaa ata gaa agc tat tgc agt aat ctc atg aag ttt gat att ccc Asp Glu Ile Glu Ser Tyr Cys Ser Asn Leu Met Lys Phe Asp Ile Pro 340 345 350	1056
gat ttc ttg gac gtt aat ggt ttt att ata taa Asp Phe Leu Asp Val Asn Gly Phe Ile Ile 355 360	1089

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MBI-17 Sequence Listing ST25

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Trp Thr Ser Glu Glu Asp Gln Lys Leu Val Asp Tyr Ile Gln Lys His
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Gly Tyr Gly Asn Trp Arg Thr Leu Pro Lys Asn Ala Gly Thr Cys Leu
35 40 45

Gln Arg Cys Gly Lys Ser Cys Arg Leu Arg Trp Thr Asn Tyr Leu Arg
50 55 60

Pro Asp Ile Lys Arg Gly Arg Phe Ser Phe Glu Glu Glu Ala Ile
65 70 75 80

Ile Gln Leu His Ser Phe Leu Gly Asn Lys Trp Ser Ala Ile Ala Ala
85 90 95

Arg Leu Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Phe Trp Asn Thr
100 105 110

His Ile Arg Lys Lys Leu Leu Arg Met Gly Ile Asp Pro Val Thr His
115 120 125

Ser Pro Arg Leu Asp Leu Leu Asp Ile Ser Ser Ile Leu Ala Ser Ser
130 135 140

Leu Tyr Asn Ser Ser His His Met Asn Met Ser Arg Leu Met Met
145 150 155 160

Asp Thr Asn Arg Arg His Gln Gln Gln His Pro Leu Val Asn Pro Glu
165 170 175

Ile Leu Lys Leu Ala Thr Ser Ile Phe Ser Gln Asn Gln Asn Asn
180 185 190

His Asn Gln Asn Gln Asn Gln Asn Gln Asn Leu Val Val Asp His Glu
195 200 205

Lys Gln Thr Val Tyr His His Asp Val Asn Gln Thr Gly Val Asn
210 215 220

Gln Tyr Gln Thr Asp Gln Tyr Phe Glu Asn Ala Ile Thr Gln Glu Leu
225 230 235 240

Gln Ser Ser Met Pro Pro Phe Pro Asn Glu Ala His Gln Phe Asn Asp
245 250 255

Met Asp His His Phe Asn Gly Phe Glu Gln Asn Leu Val Ser Thr
260 265 270

Ser Thr Thr Ser Val Gln Asp Cys Tyr Asn Pro Ser Phe Asn Asp Tyr
275 280 285

MBI-17 Sequence Listing.ST25
Ser Ser Ser Asn Phe Val Leu Asp His Ser Tyr Ser Asp Gln Ser Phe
290 295 300

Asn Phe Ala Asn Ser Val Leu Asn Thr Pro Ser Ser Ser Pro Ser Pro
305 310 315 320

Thr Thr Leu Asn Ser Ser Tyr Ile Asn Ser Ser Ser Cys Ser Thr Glu
325 330 335

Asp Glu Ile Glu Ser Tyr Cys Ser Asn Leu Met Lys Phe Asp Ile Pro
340 345 350

Asp Phe Leu Asp Val Asn Gly Phe Ile Ile
355 360

INTERNATIONAL SEARCH REPORT

Intern.....al application No.

PCT/US00/31457

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A01H 1/00, 5/00; A61K 38/10; C07H 21/00; C12N 5/14, 15/11, 15/29, 15/82
 US CL : 435/468,419,320.1;530/300,326,327;536/23.1,23.6;800/278,281,287,305-310,314,315,317.1-317.4,320.1-320.3

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/468,419,320.1;530/300,326,327;536/23.1,23.6;800/278,281,287,305-310,314,315,317.1-317.4,320.1-320.3

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
EAST, STN (Agricola, Biosis, Caplus, Embase), SEQ ID NO: 1&2**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	LI, S.F. et al. A novel myb-related gene from <i>Arabidopsis thaliana</i> . FEBS Letters 1996, Vol. 379, pages 117-121, entire reference	1-14, 25 & 26
Y		----- 27
X	SCHAFFER, R. et al. The late elongated hypocotyl mutation of <i>Arabidopsis</i> disrupts circadian rhythms and the photoperiodic control of flowering. Cell 1998, Vol. 93, pages 1219-1229.	1-14, 25 & 26
Y	Database NCBI Nucleotide, U.S. National Library of Medicine, (Bethesda, MD, USA), No. U28422, WANG, Z. Direct Submission, Sequence, January 14, 1997.	1-14 & 25-27
Y	US 5,939,601 (KLESSIG et al) 17 August 1999 (17.08.1999), entire reference.	1-14 & 25-27
Y	SUZUKI, A. et al. Cloning and expression of five myb-related genes from rice seed. Gene 1997, Vol. 198, pages 393-398.	1-14 & 25-27
Y,P	LOGUERCIO, L.L. et al. Differential regulation of six novel myb-domain genes defines two distinct expression patterns in allotetraploid cotton (<i>Gossypium hirsutum</i> L.), Mol. Gen. Genet. 1999, Vol. 261, pages 660-671.	1-14 & 25-27
Y,P	KIRIK, V. et al. Two novel myb homologues with changed expression in late embryogenesis-defective <i>Arabidopsis</i> mutants. Plant Mol. Biol. 1998, Vol. 37, pages 819-827.	1-14 & 25-27

 Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T"

later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X"

document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y"

document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&"

document member of the same patent family

Date of the actual completion of the international search

14 February 2001 (14.02.2001)

Date of mailing of the international search report

19 MAR 2001

Name and mailing address of the ISA/US
 Commissioner of Patents and Trademarks
 Box PCT
 Washington, D.C. 20231
 Facsimile No. (703)305-3230

Authorized officer

David Kruse

Telephone No. 703-308-0196

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US00/31457

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claim Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

2. Claim Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. Claim Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:
Please See Continuation Sheet

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 1-14 &25-27:SEQ ID NOS: 1&2

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US00/31457

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Groups I-XXVII, claim(s) 1-14 and 25-27, drawn to a transgenic plant having modified seed characteristics, polynucleotides and vectors for producing said transgenic plant and a method of making said transgenic plant. Applicant must elect one pair of sequences (one nucleic acid and the corresponding amino acid translation) to be examined, i.e. SEQ ID NO: 1 and 2 in Group I, SEQ ID NO: 3 and 4 in Group II, SEQ ID NO: 5 and 6 in Group III, etc.

Group XXVIII, claim(s) 15-17, drawn to a method of identifying a factor that is modulated.

Group XXIX, claims(s) 18, drawn to a method of identifying a molecule that modulates activity or expression of a polynucleotide or polypeptide.

Group XXX, claims(s) 19 and 20, drawn to an integrated computer system.

Group XXXI, claim(s) 21-24, drawn to a method for identifying a polynucleotide sequence comprising selecting a nucleic acid sequence from a database that meets a selected sequence criteria.

The inventions listed as Groups I-XXXI do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The inventions listed as Groups I-XXXI do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: Groups I-XXVII are drawn to a transgenic plant and a method of producing said plant with a nucleic acid sequence. The methods of Groups I-XXVII differ from each other in that they are directed to a plant transformation method and transgenic plant with a structurally and functionally distinct nucleic acid sequence which encodes a structurally and functionally distinct amino acid sequence. In addition, Groups XXVIII, XXIX and XXXI are different methods from any of Groups I-XXVII in that they have different method steps and different end products, and Group XXX requires a computer system. Thus, there is no single special technical feature, which links the inventions of Groups I-XXXI under PCT Rule 13.2.